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**WO 01/81405 A2**

(54) Title: METHODS AND COMPOSITIONS FOR THE PREVENTION AND TREATMENT OF ANEMIA

(57) Abstract: Methods for increasing and maintaining hematocrit in a mammal comprising administering a hyperglycosylated analog of erythropoietin are disclosed. An analog may be administered less frequently than an equivalent molar amount of recombinant human erythropoietin to obtain a comparable target hematocrit and treat anemia. Alternatively, a lower molar amount of a hyperglycosylated analog may be administered to obtain a comparable target hematocrit and treat anemia. Also disclosed are new hyperglycosylated erythropoietin analogs, methods of production of the analogs, and compositions comprising the analogs.

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**METHODS AND COMPOSITIONS FOR THE PREVENTION AND  
TREATMENT OF ANEMIA**

5    Field of the Invention

The invention relates to increasing hematocrit in a mammal using hyperglycosylated analogs of erythropoietin. More particularly, the invention relates to less frequent dosing of a hyperglycosylated 10 analog compared to recombinant human erythropoietin to raise and maintain hematocrit and treat anemia. The invention also relates to administration of lower amounts of a hyperglycosylated analog compared to recombinant human erythropoietin at an equivalent 15 dosing frequency in order to raise and maintain hematocrit and treat anemia. New hyperglycosylated analogs of erythropoietin are also provided.

Background of the Invention

20            Erythropoietin (Epo) is a glycoprotein hormone necessary for the maturation of erythroid progenitor cells into erythrocytes. It is produced in the kidney and is essential in regulating levels of red blood cells in the circulation. Conditions marked by 25 low levels of tissue oxygen signal increased production of Epo, which in turn stimulates erythropoiesis. A loss of kidney function as is seen in chronic renal failure (CRF), for example, typically results in decreased production of Epo and a concomitant reduction 30 in red blood cells.

          Human urinary Epo was purified by Miyake et al. (J. Biol. Chem. 252, 5558 (1977)) from patients with aplastic anemia. However, the amount of purified Epo protein obtained from this source was insufficient 35 for therapeutic applications. The identification and

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cloning of the gene encoding human Epo and expression of recombinant protein was disclosed in U.S. Patent No. 4,703,008 to Lin, the disclosure of which is incorporated herein by reference. A method for 5 purification of recombinant human erythropoietin from cell medium is disclosed in U.S. Patent No. 4,667,016 to Lai et. al., which is incorporated herein by reference. The production of biologically active Epo from mammalian host cells has made available, for the 10 first time, quantities of Epo suitable for therapeutic applications. In addition, knowledge of the gene sequence and the increased availability of purified protein has led to a better understanding of the mode of action of this protein.

15 Both human urinary derived Epo (Miyake et al. supra) and recombinant human Epo expressed in mammalian cells contain three N-linked and one O-linked oligosaccharide chains which together comprise about 40% of the total molecular weight of the glycoprotein. 20 N-linked glycosylation occurs at asparagine residues located at positions 24, 38 and 83 while O-linked glycosylation occurs at a serine residue located at position 126 (Lai et al. *J. Biol. Chem.* 261, 3116 (1986); Broudy et al. *Arch. Biochem. Biophys.* 265, 329 25 (1988)). The oligosaccharide chains have been shown to be modified with terminal sialic acid residues with N-linked chains typically having up to four sialic acids per chain and O-linked chains having up to two sialic acids. An Epo polypeptide may therefore 30 accommodate up to a total of 14 sialic acids.

35 Various studies have shown that alterations of Epo carbohydrate chains can affect biological activity. In one study, however, the removal of N-linked or O-linked oligosaccharide chains singly or together by mutagenesis of asparagine or serine

residues that are glycosylation sites sharply reduces in vitro activity of the altered Epo that is produced in mammalian cells (Dube et. al. J. Biol. Chem. 263, 17516 (1988)). However, DeLorme et al. (Biochemistry 31, 9871-9876 (1992)) reported that removal of N-linked glycosylation sites in Epo reduced in vivo but not in vitro biological activity.

The relationship between the sialic acid content of Epo and in vivo biological activity was disclosed by determining the in vivo activity of isolated Epo isoforms. It was found that a stepwise increase in sialic acid content per Epo molecule gave a corresponding stepwise increase in in vivo biological activity as measured by the ability of equimolar concentrations of isolated Epo isoforms to raise the hematocrit of normal mice (Egrie et al. Glycoconjugate J. 10, 263 (1993)). Those Epo isoforms having higher sialic acid content also exhibited a longer serum half-life but decreased affinity for the Epo receptor, suggesting that serum half-life is an important determinant of in vivo biological activity.

Introduction of new glycosylation sites in the Epo polypeptide can result in the production of molecules with additional carbohydrate chains. See PCT Publication Nos. WO91/05867 and WO94/09257 hereby incorporated by reference in their entirety. Epo glycosylation analogs having at least one additional N-linked carbohydrate chain and/or having at least one additional O-linked carbohydrate chain are disclosed. A glycosylation analog having one additional N-linked chain was determined to have a longer circulating half-life compared to recombinant human Epo (rHuEpo) (isoforms 9-14) and to a purified isoform of rHuEpo having 14 sialic acids per molecule.

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Administration of recombinant human erythropoietin (rHuEpo) is effective in raising red blood cell levels in anemic patients with end stage renal disease (Eschbach et al. *New Eng. J. Med.* 316, 5 73-38 (1987)). Subsequent studies have shown that treatment with rHuEpo can correct anemia associated with a variety of other conditions. (Fischl et al. *New Eng. J. Med.* 322, 1488-1493 (1990); Laupacis, *Lancet* 341, 1228-1232 (1993)). Regulatory approvals have been 10 given for the use of rHuEpo in the treatment of anemia associated with CRF, anemia related to therapy with AZT (zidovudine) in HIV-infected patients, anemia in patients with non-myeloid malignancies receiving chemotherapy, and anemia in patients undergoing surgery 15 to reduce the need of allogenic blood transfusions. Current therapy for all approved indications (except the surgery indication) involves a starting dose of between 50-150 Units/kg three times per week (TIW) administered either by an intravenous (IV) or 20 subcutaneous (SC) injection to reach a suggested target hematocrit range. For the surgery indication, rHuEpo is administered every day 10 days prior to surgery, on the day of surgery, and four days thereafter (EPOGEN® Package Insert, 12/23/96). In general, the current 25 recommended starting doses for rHuEpo raise hematocrit into the target range in about six to eight weeks. Once the target hematocrit range has been achieved, a maintenance dosing schedule is established which will vary depending upon the patient, but is typically three 30 times per week for anemic patients with CRF. The administration of rHuEpo described above is an effective and well-tolerated regimen for the treatment of anemia.

It would be desirable to have a therapeutic 35 with greater potency than rHuEpo. An advantage to such

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a molecule would be that it could be administered less frequently and/or at a lower dose. Current treatments for patients suffering from anemia call for administration of EPOGEN® three times per week and for 5 surgery patients administration once per day. A less frequent dosing schedule would be more convenient to both physicians and patients, especially those patients who do not make regularly scheduled visits to doctor's offices or clinics, or those who self-inject their Epo. 10 Another advantage of a more potent molecule is that less drug is being introduced into patients for a comparable increase in hematocrit.

It is therefore an object of the invention to identify more potent molecules for the treatment of 15 anemia which will permit a less frequent dosing schedule. It is a further object of the invention to provide molecules which will increase and maintain hematocrit at levels which are at least comparable to that of Epo when administered at a lower dose. It is 20 also an object of the invention that these molecules selected for less frequent dosing is at least as well tolerated as rHuEpo and potentially better tolerated in some patients.

25 Summary of the Invention

It has been found that a hyperglycosylated Epo analog designated N47 (Asn<sup>30</sup>Thr<sup>32</sup>Val<sup>87</sup>Asn<sup>88</sup>Thr<sup>90</sup> Epo) has a longer serum half-life than recombinant human erythropoietin (rHuEpo) and a greater in vivo activity 30 when administered at the same dose and frequency as rHuEpo. Further, the analog has been shown to raise hematocrit in mice at once per week administration that is comparable to hematocrit rise for rHuEpo administered three times per week. The

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pharmacokinetics of Epo analog N47 administered to mice and to humans were similar.

The invention provides for a method of raising and maintaining hematocrit in a mammal comprising administering a therapeutically effective amount of an Epo hyperglycosylated analog in a pharmaceutical composition, wherein the analog is administered less frequently than an equivalent molar amount of rHuEpo to obtain a comparable target hematocrit. The dosing frequency of the present invention in order to reach a patient's optimal hematocrit range is less than three times per week. Dosing frequencies may be two times per week, one time per week, or less than one time per week, such as one time every other week, once per month or once every two months. The dosing frequency required to maintain a patient's target hematocrit is less than three times per week. Dosing frequencies may be two times per week, one time per week, or less than one time per week, such as one time every two weeks, once per month or once every two months.

The invention also provides for a method of raising and maintaining hematocrit in a mammal comprising administrating a therapeutically effective amount of an Epo hyperglycosylated analog wherein the analog is administered at a lower molar amount than rHuEpo to obtain a comparable target hematocrit.

Also provided for are pharmaceutical compositions comprising Epo hyperglycosylated analogs wherein the compositions are suitable for dosing frequency of less than three times per week. The compositions will include pharmaceutically acceptable adjuvants suitable for use with Epo hyperglycosylated analogs.

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The invention may be employed with any condition resulting in a decrease in red blood cell levels, such as anemia associated with a decline or loss of kidney function, (chronic renal failure) 5 myelosuppressive therapy, cancer, viral infection, chronic disease and excessive loss of blood during surgical procedures. In one embodiment, treatment with once per week dosing, or less frequently, is for anemia resulting from chronic renal failure.

10 Also provided for are new hyperglycosylated analogs of Epo. The analogs comprise at least one additional carbohydrate chain compared to rHuEpo wherein at least one N-linked carbohydrate chain is added at any of positions 52, 53, 55, 86 and 114. New 15 hyperglycosylated analogs may have two, three or four additional carbohydrate chains, or may have more than four additional chains.

Description of the Figures

20 Figure 1 (SEQ ID NO: 1) shows the amino acid sequence of human erythropoietin.

25 Figure 2 shows a Western blot analysis of rHuEpo and Epo hyperglycosylated analogs from CHO cell expression in serum free medium. Construction of analogs N53 and N61 are described in Example 1. The number of N-linked carbohydrate chains on each analog is indicated.

30 Figure 3 compares the activity of rHuEpo, Epo analogs N4, N18, and N50 (containing four N-linked carbohydrate chains), N47 (containing five N-linked carbohydrate chains), and N53 (containing six N-linked carbohydrate chains) in the exhypoxic polycythemic 35 mouse bioassay. Experimental procedures are described

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in Example 3. Each point represents the mean response of five animals. Analogs N4, N18 and N47 have been described previously in WO94/09257.

5                 Figure 4 compares the serum half-life of rHuEpo and Epo analog N47 administered to normal rats by intravenous injection (IV). Experimental procedures are described in Example 4. Results are the mean ( $\pm$ SD) for each group.

10                 Figure 5 compares the serum half-life of rHuEpo and Epo analog N47 administered to Beagle dogs by intravenous injection (IV). Experimental procedures are described in Example 4. Results are the mean ( $\pm$ SD) 15 for each group.

20                 Figure 6 shows the increase in hematocrit in mice in response to varying doses of rHuEpo or Epo analog N47 administered by intraperitoneal injection (IP) three times per week (TIW) for six weeks. Experimental procedures are described in Example 5. Results shown are the group mean ( $\pm$ SD) of the change in hematocrit for each dose group.

25                 Figure 7 compares the relative potency in mice of rHuEpo and Epo analog N47 injected by either the intraperitoneal (IP) or intravenous (IV) routes of administration at a frequency of once weekly (QW) or three time a week (TIW). Experimental procedures are 30 described in Example 5. Each point represents the mean ( $\pm$ SD) of data from separate experiments as follows: N47, IP, TIW (n=5); N47, IV, TIW (n=1); N47, IP, QW (n=2); N47, IV, QW (n=3); rHuEpo, IP, TIW (n=5); rHuEpo, IV, QW (n=2). Each experiment used 7 - 13 mice 35 per dose.

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Figure 8 shows the increase in hematocrit in mice in response to varying doses of rHuEpo or Epo analog N47 administered by intravenous (IV) injection 5 one time per week (QW) for approximately six weeks. Experimental procedures are described in Example 5. Results shown are the group mean ( $\pm$ SD) of the change in hematocrit for each dose group.

10 Figure 9 shows the increase in hematocrit in mice in response to varying doses of Epo analog N47 administered by intravenous (IV) injection one time per week (QW) or once every other week (EOW) for approximately six weeks. Experimental procedures are 15 described in Example 5. Results shown are the group mean ( $\pm$ SD) of the change in hematocrit for each dose group.

20 Figure 10 (SEQ ID NO: 25) shows the amino acid sequence of the hinge, CH<sub>2</sub> and CH<sub>3</sub> regions of human IgG<sub>1</sub>.

25 Figure 11 (SEQ ID NO: 26) shows the cDNA and amino acid sequence of Epo N47-Fc fusion polypeptide including the Epo signal sequence. The amino terminal Fc residue is fused to the arg-166 residue of Epo.

#### Detailed Description of the Invention

30 The invention provides for a method of raising and maintaining hematocrit comprising administering a therapeutically effective amount of a hyperglycosylated analog of erythropoietin in a pharmaceutical composition. The analog is administered less frequently than an equivalent molar amount of 35 rHuEpo to obtain a comparable target hematocrit. The

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invention also provides for a method of raising and maintaining hematocrit comprising administering a hyperglycosylated analog in lower molar amounts than rHuEpo to obtain a comparable target hematocrit. The 5 composition may be administered by intravenous, subcutaneous or intraperitoneal routes.

Surprisingly, it has been found that analog N47, a hyperglycosylated Epo analog described in WO94/09257, could achieve an increase in hematocrit 10 administered once a week that was comparable to that observed for rHuEpo given three times per week. Analog N47 has the following amino acid changes: ala to asn at 30; his to thr at 32; pro to val at 87; trp to asn at 88; and pro to thr at 90 which resulted in the addition 15 of two N-linked carbohydrate chains at asparagine residues 30 and 88. The analog was expressed in Chinese hamster ovary (CHO) cells (as described in Example 1) and purified as described in Example 2 to give isoforms of 17 to 22 sialic acids. Analog N47 20 showed a greater serum half-life in rats and beagle dogs than rHuEpo when injected intravenously (Figures 4 and 5). When injected intraperitoneally three times per week, N47 induced increases in hematocrit of normal mice comparable to rHuEpo at lower concentrations 25 (Figure 6). The potency of N47 was demonstrated to be about 3 to 4-fold higher than rHuEpo when administered three times per week (Figures 6 and 7). When given once per week, at similar doses, rHuEpo showed little stimulation of hematocrit in normal mice while N47 gave 30 a marked increase (Figure 8). The potency of N47 was about 14-fold higher than rHuEpo for once per week dosing (Figure 7). Significantly, the hematocrit response for analog N47 given once per week is comparable to that for rHuEpo given three times per 35 week. Even when administered once every other week,

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N47 still produced significant increases in the hematocrit of normal mice (Figure 9). Taken together, the data indicated that Epo hyperglycosylated analogs, and analog N47 in particular, can be used 5 advantageously to raise hematocrit using less frequent dosing than for current treatment with rHuEpo.

It has also been shown that the results described above obtained in mice may be extrapolated to humans. Pharmacokinetic parameters for administration 10 of rHuEpo and analog N47 to 11 Continuous Ambulatory Peritoneal Dialysis (CAPD) patients demonstrate that analog N47 has a three-fold longer serum half-life than rHuEpo (Example 6 and Table 5). These results suggest that Epo hyperglycosylated analogs allow less frequent 15 dosing than rHuEpo in humans.

As used herein, the term "hyperglycosylated Epo analog" refers to Epo comprising at least one additional glycosylation site with an additional carbohydrate chain added to the site. Glycosylation 20 sites may be for N-linked or O-linked carbohydrate chains. New N-linked glycosylation sites are introduced by alterations in the DNA sequence to encode the consensus site for N-linked carbohydrate addition (the amino acids Asn-X-Ser/Thr) in the polypeptide 25 chain, while new O-linked sites are introduced by alterations in the DNA sequence to encode a serine or a threonine residue. The analogs are constructed by mutagenesis techniques for introducing additions, deletions or substitutions of amino acid residues that 30 increase or alter sites in the Epo polypeptide that are available for glycosylation. DNA encoding an Epo hyperglycosylated analog is transfected into a eucaryotic host cell and the expressed glycoprotein is analyzed for the presence of an additional carbohydrate 35 chain.

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Epo hyperglycosylated analogs have shown in vitro activity which was comparable to or even less than that determined for rHuEpo, suggesting that binding to the Epo receptor is not enhanced, and may in 5 some cases be diminished, by addition of carbohydrate chains. However, hyperglycosylation can typically increase serum half-life and potentially lead to increased in vivo biological activity. One Epo analog having an additional N-linked carbohydrate chain at 10 position 88 exhibited decreased affinity for receptor compared to rHuEpo (isoforms 9-14) or to a purified isoform of rHuEpo having 14 sialic acids per molecule, yet demonstrated a longer circulating half-life and enhanced in vivo activity compared to either a mixture 15 of Epo isoforms 9-14 or isolated Epo isoform 14.

The Epo hyperglycosylated analogs which may be administered according to the present invention will have at least one additional N-linked or O-linked carbohydrate chain. In one embodiment, the analogs 20 will have two additional N-linked carbohydrate chains. In other embodiments, the analogs will have three, four or more additional N-linked carbohydrate chains. As examples, the analogs of the invention will have at least one additional N-linked chain at one or more of 25 amino acid residues 30, 51, 57, 69, 88, 89, 136 and 138 of the sequence of human Epo. In one embodiment, the analog has additional N-linked carbohydrate chains at residues 30 and 88 of human Epo. The numbering of amino acid residues of human Epo is as shown in Figure 30 1 and SEQ ID NO:1. Figure 1 shows a predicted mature Epo polypeptide of 166 amino acids whereas recombinant produced Epo has 165 amino acids after removal of the C-terminal arginine residue. It is understood that rHuEpo and hyperglycosylated Epo analogs may have 35 either 165 or 166 amino acids.

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The analogs of the invention will have at least four N-linked carbohydrate chains. Of the four chains, three may be at the naturally occurring sites at positions 24, 38, and 83. However, it is 5 contemplated that some analogs of the invention may have alterations of one or more of the naturally occurring glycosylation sites such that one or more of the sites are deleted and substituted with a new site. Such analogs are also provided by the invention. For 10 example, any one of sites at positions 24, 38 and 83 may be deleted and substituted with a site at position 88. Optionally, the analogs may have an O-linked site at position 126.

The invention also provides for new Epo 15 hyperglycosylated analogs having at least one additional carbohydrate chain. It has been found that an additional N-linked carbohydrate chain is added at any of positions 52, 53, 55, 86 and 114 which have been modified to be a glycosylation site. Specific 20 embodiments include analogs N49 through N61 as described in Table 1. The new analogs will have at least one new N-linked glycosylation site at any of positions 52, 53, 55, 86 and 114 and may further comprise additional N-linked or O-linked carbohydrate 25 chains at other sites. The analogs may have one, two, three or four additional carbohydrate chains, or more than four additional chains. In one preferred embodiment, the analogs will have three additional N-linked carbohydrate chains (six N-linked chains total). In another preferred embodiment, the analogs 30 will have four additional N-linked chains (seven N-linked chains total). The analogs having three or four, or more than four, additional N-linked carbohydrate chains may have, but are not limited to,

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an additional chain at any of positions 52, 53, 55, 86 and 114.

Surprisingly, it has been found that a hyperglycosylated analog with three additional N-linked chains at positions 30, 53 and 88 (six N-linked chains total) has a greater in vivo activity than analog N47 with two additional chains (five total). The results are shown in Figure 3. It is clear that the in vivo activity of the analogs is directly dependent on the number of N-linked carbohydrate chains. These results may be extrapolated to the therapeutic setting wherein the analogs having more N-linked carbohydrate chains than N47 may be dosed even less frequently.

In addition, the invention provides for hyperglycosylated Epo analogs with three additional N-linked chains at positions 30, 55 and 88; 30, 55 and 114; and 30, 88 and 114. Epo analogs with four additional N-linked chains or three additional N-linked chains and one additional O-linked chain at position 125 are also provided.

The analogs may be prepared by a variety of mutagenesis techniques available to one skilled in the art, such as site-directed mutagenesis, PCR mutagenesis and cassette mutagenesis (Zoller et al. *Meth. Enz.* 100, 468-500 (1983); Higuchi, in PCR Protocols pp. 177-183 (Academic Press, 1990); Wells et al. *Gene* 34, 315-323 (1985)). Example 1 describes the use of PCR mutagenesis techniques to construct new Epo hyperglycosylated analogs.

An Epo DNA sequence which has undergone mutagenesis is inserted into an expression vector using standard techniques with the vector being suitable for maintenance in a mammalian host cell. The vector will typically contain the following elements: promoter and other "upstream" regulatory elements, origin of

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replication, ribosome binding site, transcription termination site, polylinker site, and selectable marker that are compatible with use in a mammalian host cell. Vectors may also contain elements that allow 5 propagation and maintenance in prokaryotic host cells as well.

Suitable cells or cell lines include any from mammalian sources, including human sources. Examples include COS-7 (ATCC accession no. CRL 1651), human 293, 10 baby hamster kidney (BHK, ATCC accession no. CCL 10), Chinese hamster ovary cells (including dihydrofolate reductase (DHFR)-deficient cells, Urlab et al. Proc. Natl. Acad. Sci. USA 77, 4216-4220 (1980)) Other suitable mammalian cell lines include, but are not 15 limited to, HeLa, mouse L-929 and 3T3. In a preferred embodiment, DHFR-deficient CHO cells are used.

Vectors comprising sequences encoding Epo hyperglycosylation analogs are introduced into host cells by standard transformation or transfection 20 techniques. Culturing, amplifying and screening transformed or transfected host cells are accomplished using publicly available methods (Gething et al. Nature 293, 620-625 (1981); Kaufman et al. Mol Cell. Biol. 5, 1750-1759 (1985); U.S. Patent No. 4,419,446). Host 25 cells harboring DNA sequences encoding Epo hyperglycosylated analogs are cultured under conditions that permit expression of the analogs. The analogs are recovered from the cell media and purified using procedures essentially as described previously 30 (WO94/09257) and those in Example 2. The purification procedures allow for the isolation of higher sialic acid containing Epo isoforms resulting from adding additional carbohydrate chains.

The Epo hyperglycosylated analogs may 35 include, in addition to new glycosylation sites,

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additions, deletions or substitutions of amino acid residues which do not create new glycosylation sites and do not substantially alter the biological activity of the hyperglycosylated analog. Those individual 5 sites or regions of Epo which may be altered without affecting biological activity may be determined by examination of the structure of the Epo-Epo receptor complex as described in Syed et. al. *Nature* 395, 511 (1998). Examination of the structure of the Epo-Epo 10 receptor complex reveals those residues which interact with, or are in close proximity to, the receptor binding site of Epo and which should be avoided when making alterations in the Epo amino acid sequence. Alternatively, one may empirically determine those 15 regions which would tolerate amino acid substitutions by alanine scanning mutagenesis (Cunningham et al. *Science* 244, 1081-1085 (1989). In this method, selected amino acid residues are individually substituted with a neutral amino acid (e.g., alanine) 20 in order to determine the effects on biological activity.

It is generally recognized that conservative amino acid changes are least likely to perturb the structure and/or function of a polypeptide.

25 Accordingly, the invention encompasses one or more conservative amino acid changes within an Epo hyperglycosylated analog. Conservative amino acid changes generally involve substitution of one amino acid with another that is similar in structure and/or 30 function (e.g., amino acids with side chains similar in size, charge and shape). The nature of these changes are well known to one skilled in the art and are summarized in Table 1 below. Such conservative substitutions are shown under the heading of "Preferred 35 substitutions". Also contemplated are more substantial

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changes ("Exemplary substitutions") which may also be introduced. A skilled artisan will appreciate that initially the sites should be modified by substitution in a relatively conservative manner. If such 5 substitutions result in a retention in biological activity, then more substantial changes (Exemplary Substitutions) may be introduced and/or other additions/deletions may be made and the resulting products screened.

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TABLE 1: Amino Acid Substitutions

<u>Original Residue</u>	<u>Preferred Substitutions</u>	<u>Exemplary Substitutions</u>
Ala (A)	Val	Val; Leu; Ile
Arg (R)	Lys	Lys; Gln; Asn
Asn (N)	Gln	Gln; His; Lys; Arg
Asp (D)	Glu	Glu
Cys (C)	Ser	Ser
Gln (Q)	Asn	Asn
Glu (E)	Asp	Asp
Gly (G)	Pro	Pro
His (H)	Arg	Asn; Gln; Lys; Arg
Ile (I)	Leu	Leu; Val; Met; Ala; Phe; norleucine
Leu (L)	Ile	norleucine; Ile; Val; Met; Ala; Phe
Lys (K)	Arg	Arg; Gln; Asn
Met (M)	Leu	Leu; Phe; Ile
Phe (F)	Leu	Leu; Val; Ile; Ala
Pro (P)	Gly	Gly
Ser (S)	Thr	Thr
Thr (T)	Ser	Ser
Trp (W)	Tyr	Tyr
Tyr (Y)	Phe	Trp; Phe; Thr; Ser
Val (V)	Leu	Ile; Leu; Met; Phe; Ala; norleucine

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Also provided by the invention are deletions or additions of amino acids in a hyperglycosylated Epo analog which do not substantially affect biological activity. Such additions and deletions may be at the 5 N-terminal or C-terminal of the polypeptide, or may be internal to it. In general, relatively small deletions or additions are less likely to affect structure and/or function of Epo or a hyperglycosylated analog. In one embodiment, deletions or additions can be from 5-10 10 residues, alternatively from 2-5 amino acid residues, or from 1-2 residues.

The invention provides for fusion proteins comprising Epo hyperglycosylated analogs and compositions thereof. In one aspect, the invention 15 provides for fusion proteins of Epo hyperglycosylated analogs and an immunoglobulin heavy chain constant region. Fusions may be made at the amino terminus of an Epo hyperglycosylated analog, that is, the carboxy terminus of an immunoglobulin heavy chain constant region. 20 The amino terminus of an Epo hyperglycosylated analog is fused to the amino terminus of an Epo hyperglycosylated analog. Alternatively, it may be desirable to fuse the carboxy terminus of an Epo hyperglycosylated analog to the amino terminus of an immunoglobulin heavy chain constant region. In one 25 aspect of the invention, the immunoglobulin heavy chain constant region is an Fc region. Epo hyperglycosylated analogs, when part of a fusion polypeptide, may be 165 or 166 amino acids in length, or may have greater or fewer residues if amino acids are added or deleted. In 30 one embodiment, analog N47 is fused at its C-terminus to the N-terminus of an Fc region derived from human IgG1. (See Example 2.) In the present example, analog N47 includes the arginine residue at position 166. However, it is contemplated that analog N47 as 35 well as other hyperglycosylated analogs of residues 1-

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165 (lacking the C-terminal arginine residue) may also comprise the fusion polypeptides of the invention.

The term "Fc" refers to a molecule or sequence comprising the sequence of a non-antigen-binding portion of antibody, whether in monomeric or multimeric form. The original immunoglobulin source of an Fc is preferably of human origin and may be from any isotype, e.g., IgG, IgA, IgM, IgE or IgD. One method of preparation of an isolated Fc molecule involves digestion of an antibody with papain to separate antigen and non-antigen binding portions of the antibody. Another method of preparation of an isolated Fc molecules is production by recombinant DNA expression followed by purification of the Fc molecules so expressed. A full-length Fc consists of the following Ig heavy chain regions: CH1, CH2 and CH3 wherein the CH1 and CH2 regions are typically connected by a flexible hinge region. In one embodiment, an Fc has the amino acid sequence of IgG1 such as that shown in Figure 10. The terms "Fc protein", "Fc sequence", "Fc molecule", "Fc region" and "Fc portion" are taken to have the same meaning as "Fc".

The term "Fc fragment" when used in association with Fc molecule, or fusion polypeptides thereof, refers to a peptide or polypeptide that comprises less than the full length amino acid sequence of an Fc molecule. Such a fragment may arise, for example, from a truncation at the amino terminus, a truncation at the carboxy terminus, and/or an internal deletion of a residue(s) from the amino acid sequence. Fc fragments may result from alternative RNA splicing or from *in vivo* protease activity.

The term "Fc variant" when used in association with an Fc molecule, or with fusion

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polypeptides thereof, refers to a polypeptide comprising an amino acid sequence which contain one or more amino acid sequence substitutions, deletions, and/or additions as compared to native Fc amino acid sequences. Variants may be naturally occurring or artificially constructed. Variants of the invention may be prepared from the corresponding nucleic acid molecules encoding said variants, which have a DNA sequence that varies accordingly from the DNA sequences for native Fc molecule.

The term "derivative" when used in association with an Fc molecule, or with fusion polypeptides thereof, refers to Fc variants or fragments thereof, that have been chemically modified, as for example, by covalent attachment of one or more polymers, including, but limited to, water soluble polymers, N-linked or O-linked carbohydrates, sugars, phosphates, and/or other such molecules. The derivatives are modified in a manner that is different from native Fc, either in the type or location of the molecules attached to the polypeptide. Derivatives further includes deletion of one or more chemical groups naturally attached to an Fc molecule.

The term "fusion" refers to joining of different peptide or protein segments by genetic or chemical methods wherein the joined ends of the peptide or protein segments may be directly adjacent to each other or may be separated by linker or spacer moieties such as amino acid residues or other linking groups.

An Fc, or a variant, fragment or derivative thereof, may be from an Ig class. In one embodiment, an Fc is from the IgG class, such as IgG1, IgG2, IgG3, and IgG4. In another embodiment, an Fc is from IgG1. An Fc may also comprise amino acid residues represented by a combination of any two or more of the Ig classes,

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such as residues from IgG1 and IgG2, or from IgG1, IgG2 and IgG3, and so forth. In one embodiment, an Fc region of an Epo hyperglycosylated analog fusion protein has the sequence as set forth in Figure 10 (SEQ 5 ID NO: 25) (see Ellison et al., Nucleic Acids Res. 10, 4071-4079 (1982)) starting at residue 6 (that is, residues 1-5 are deleted).

In addition to naturally occurring variations in Fc regions, Fc variants, fragments and derivatives 10 may contain non-naturally occurring changes in Fc which are constructed by, for example, introducing substitutions, additions, insertions or deletions of residues or sequences in a native or naturally occurring Fc, or by modifying the Fc portion by 15 chemical modification and the like. In general, Fc variants, fragments and derivatives are prepared such that the increased circulating half-life of Fc fusions to Epo glycosylation analogs is largely retained.

Also provided by the invention are Fc 20 variants with conservative amino acid substitutions. Examples of conservative amino acid substitutions are set forth hereinabove, and are also exemplified by substitution of non-naturally occurring amino acid residues which are typically incorporated by chemical 25 peptide synthesis rather than by synthesis in biological systems. These include peptidomimetics, and other reversed or inverted forms of amino acid moieties. Conservative modifications to the amino acid sequence of an Fc region (and the corresponding 30 modifications to the encoding nucleotides) are expected to produce Fc molecules (and fusion proteins comprising Epo hyperglycosylated analogs and Fc regions) which have functional and chemical characteristics similar to those of unmodified Fc molecules and fusion proteins 35 comprising unmodified Fc regions.

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In addition to the substitutions set forth in Table I, any native residue in an Fc molecule (or in an Fc region of a fusion protein comprising an Epo hyperglycosylated analog) may also be substituted with 5 alanine, as has been previously described for "alanine scanning mutagenesis" (Cunningham et al. *Science* 244, 1081-1085 (1989)).

Substantial modifications in the functional and/or chemical characteristics of an Fc molecule (and 10 in an Fc region of a fusion protein comprising an Epo hyperglycosylated analog) may be accomplished by selecting substitutions that differ significantly in their effect on maintaining (a) the structure of the molecular backbone in the area of the substitution, for 15 example, as a sheet or helical conformation, (b) the charge or hydrophobicity of the molecule at the target site, or (c) the bulk of the side chain. Naturally occurring residues may be divided into groups based on common side chain properties:

- 20 1) hydrophobic: norleucine, Met, Ala, Val, Leu, Ile;
- 2) neutral hydrophilic: Cys, Ser, Thr;
- 3) acidic: Asp, Glu;
- 4) basic: Asn, Gln, His, Lys, Arg;
- 25 5) residues that influence chain orientation: Gly, Pro; and
- 6) aromatic: Trp, Tyr, Phe.

Non-conservative substitutions may involve the exchange of a member of one of these classes for a 30 member from another class. Such substituted residues may be introduced into regions of an Fc molecule that are homologous with a non-human Fc molecule, or into the non-homologous regions of the molecule.

Cysteine residues in Fc molecules can be 35 deleted or replaced with other amino acids to prevent

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formation of disulfide crosslinks. In particular, a cysteine residue at position 5 of Figure 10 (SEQ. ID. NO. \_\_) may be substituted with one or more amino acids, such as alanine or serine. Alternatively, the 5 cysteine residue at position 5 could be deleted.

An Fc fragment may be prepared by deletion of one or more amino acids at any of positions 1, 2, 3, 4 and 5 as shown in Figure 10 (SEQ ID NO. 25). In one embodiment, the amino acid residues at positions 1-5 10 inclusive are deleted. Substitutions at these positions can also be made and are within the scope of this invention.

Fc variants may also be made which show reduced binding to Fc receptors which trigger effector 15 functions such as antibody dependent cellular cytotoxicity (ADCC) and activation of complement (see for example Molec. Immunol. 29, 633-639, (1992)). Such variants may include leucine at position 20 deleted or substituted with a glutamine residue, glutamate at 20 position 103 deleted or substituted with an alanine residue, and lysines at positions 105 and 107 deleted or substituted with alanine residues (following the numbering as set forth in Figure 1). One or more of such substitutions are contemplated.

25 In one embodiment, Fc variants will exhibit stronger binding to the FcRn receptor ("salvage receptor") and a longer circulating half-life compared to native Fc such as that shown in Figure 1. Examples of such variants include amino acid substitutions at 30 one or more of residues 33, 35-42, 59, 72, 75, 77, 95-98, 101, 172-174, 215 and 220-223, wherein the substitution(s) confer tighter binding of an Fc variant to the FcRn receptor. In another embodiment, Fc variants have one or more glycosylation sites removed. 35 N-linked glycosylation sites may be removed by deletion

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or substitution of asparagine residues having attached carbohydrate chains.

Other Fc variants include one or more tyrosine residues replaced with, for example, 5 phenylalanine residues. In addition, other variant amino acid insertions, deletions and/or substitutions are also contemplated and are within the scope of the present invention. Examples include Fc variants disclosed in WO96/32478 and WO97/34630 hereby 10 incorporated by reference. Furthermore, alterations may be in the form of altered amino acids, such as peptidomimetics or D-amino acids.

The Fc protein may be also linked to the Epo glycosylation analogs by "linker" moieties comprising 15 chemical groups or amino acids of varying lengths.

Such chemical linkers are well known in the art. Amino acid linker sequences can include but are not limited to:

- (a) ala-ala-ala;
- 20 (b) ala-ala-ala-ala; (SEQ ID NO: 6)
- (c) ala-ala-ala-ala-ala; (SEQ ID NO: 7)
- (d) gly-gly;
- (e) gly-gly-gly;
- (f) gly-gly-gly-gly-gly; (SEQ ID NO: 8)
- 25 (g) gly-gly-gly-gly-gly-gly-gly; (SEQ ID NO: 9)
- (h) gly-pro-gly;
- (i) gly-gly-pro-gly-gly; (SEQ ID NO: 10) and
- 30 (j) any combination of subparts (a) through (i).

While Fc molecules are preferred as components of fusion proteins with Epo glycosylation 35 analogs, it is also contemplated that other amino acid

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sequences which bind to an FcRn receptor and confer increased in vivo half-life may also be used. Examples of such alternative molecules are described in U.S. Pat. No. 5,739,277, issued April 14, 1998 to Presta et al.

The term "molar amount" refers to an amount of a hyperglycosylated analog or rHuEpo which is based upon the molecular weight of the corresponding erythropoietin polypeptide without glycosylation. Equivalent amounts of rHuEpo and analog refer to amounts which are equal when taking into account normal variations in procedure used to determine such amounts. It is necessary to determine equivalent amounts in this manner since the molecular weight of rHuEpo and analogs will vary depending upon the number of carbohydrate chains. For rHuEpo, the molecular weight of erythropoietin polypeptide is calculated based upon amino acid residues 1-165 as shown in Figure 1 and SEQ ID NO: 1. For hyperglycosylated analogs, the molecular weights are adjusted depending upon the amino acid changes in residues 1-165 of Figure 1 and SEQ ID NO: 1.

The dosing frequency for a hyperglycosylated analog will vary depending upon the condition being treated and the target hematocrit, but in general will be less than three times per week. The dosing frequency will be about two times per week, about one time per week. The dosing frequency may also be less than about one time per week, for example about one time every two weeks (about one time per 14 days), one time per month or one time every two months. It is understood that the dosing frequencies actually used may vary somewhat from the frequencies disclosed herein due to variations in responses by different individuals

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to the Epo analogs; the term "about" is intended to reflect such variations.

As used herein, the term "therapeutically effective amount" refers to an amount of a

5 hyperglycosylated analog (or a fusion protein comprising an Epo hyperglycosylated analog and an immunoglobulin heavy chain constant region) which gives an increase in hematocrit to a target hematocrit, or to a target hematocrit range that provides benefit to a

10 patient or, alternatively, maintains a patient at a target hematocrit, or within a target hematocrit range. The amount will vary from one individual to another and will depend upon a number of factors, including the overall physical condition of the patient, severity and

15 the underlying cause of anemia and ultimate target hematocrit for the individual patient. A target hematocrit is typically at least about 30%, or in a range of 30%-38%, preferably above 38% and more preferably 40%-45%. General guidelines relating to

20 target hematocrit ranges for rHuEpo are also found in the EPOGEN® package insert dated 12/23/96 and are 30%-36%, or alternatively 32%-38% as stated therein. It is understood that such targets will vary from one individual to another such that physician discretion

25 may be appropriate in determining an actual target hematocrit for any given patient. Nonetheless, determining a target hematocrit is well within the level of skill in the art.

A therapeutically effective amount of the

30 present compositions may be readily ascertained by one skilled in the art. Example 6 sets forth a clinical protocol which has as one objective to determine a therapeutically effective amount of analog N47 in both once per week and three times per week dosing. A dose range for once per week administration is from about

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0.075 to about 4.5  $\mu$ g erythropoietin peptide per kg per dose. A dose range for three times per week administration is 0.025 to 1.5  $\mu$ g erythropoietin peptide per kg per dose. This dose range may be 5 employed with other Epo hyperglycosylated analogs, with any adjustments in the dosing range being routine to one skilled in the art.

A significant advantage to the present invention is the ability to correlate the extent of 10 hyperglycosylation either with a dose amount or with a dosing interval that would allow one to "tailor" an Epo analog to a given dose or dosing schedule. Based upon the increasing in vivo activities of Epo analogs having one, two or three additional carbohydrate chains as 15 shown in Figure 3, the treating physician can select an analog that is appropriate and convenient for the anemic condition being treated. For example, in patients who are acutely anemic and in need of a large effective dose, or in patients which require a longer- 20 lasting treatment, administration of a hyperglycosylated analog with three or four or even more additional carbohydrate chains may be preferred. For other patients who experience less severe anemia or require treatment for a relatively short time, an 25 analog with one or two additional carbohydrate chains may be preferred. The analogs of the present invention provide the physician with considerable flexibility in preventing and treating anemia that may result from a wide variety of underlying conditions.

30 The invention also provides for administration of a therapeutically effective amount of iron in order to maintain increased erythropoiesis during therapy. The amount to be given may be readily determined by one skilled in the art based upon therapy 35 with rHuEpo.

The present invention may be used to stimulate red blood cell production and prevent and treat anemia. Among the conditions treatable by the present invention include anemia associated with a 5 decline or loss of kidney function (chronic renal failure), anemia associated with myelosuppressive therapy, such as chemotherapeutic or anti-viral drugs (such as AZT), anemia associated with the progression of non-myeloid cancers, anemia associated with viral 10 infections (such as HIV), and anemia of chronic disease. Also treatable are conditions which may lead to anemia in an otherwise healthy individual, such as an anticipated loss of blood during surgery. In general, any condition treatable with rHuEpo may also 15 be treated with the Epo hyperglycosylated analogs of the invention.

The invention also provides for pharmaceutical compositions comprising a therapeutically effective amount of an Epo 20 hyperglycosylated analog, together with a pharmaceutically acceptable diluent, carrier, solubilizer, emulsifier, preservative and/or adjuvant. The invention also provides for a pharmaceutical composition comprising a therapeutically effective 25 amount of a fusion protein comprising an Epo hyperglycosylated analog and an immunoglobulin heavy chain constant region together with a pharmaceutically acceptable diluent, carrier, solubilizer, emulsifier, preservative and/or adjuvant. The composition will be 30 suitable for a dosing schedule of less than three times per week. The composition may be in a liquid or lyophilized form and comprises a diluent (Tris, citrate, acetate or phosphate buffers) having various pH values and ionic strengths, solubilizer such as 35 Tween or Polysorbate, carriers such as human serum

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albumin or gelatin, preservatives such as thimerosal, parabens, benzylalconium chloride or benzyl alcohol, antioxidants such as ascrobic acid or sodium metabisulfite, and other components such as lysine or 5 glycine. Selection of a particular composition will depend upon a number of factors, including the condition being treated, the route of administration and the pharmacokinetic parameters desired. A more extensive survey of components suitable for 10 pharmaceutical compositions is found in Remington's Pharmaceutical Sciences, 18th ed. A.R. Gennaro, ed. Mack, Easton, PA (1980). In a preferred embodiment, the Epo glycosylation analogs of the invention are formulated in liquid form in an isotonic sodium 15 chloride/sodium citrate buffered solution containing human albumin, and optionally containing benzyl alcohol as a preservative. The compositions preferably contain analogs having one, two, three, four, or more additional carbohydrate chains.

20 Compositions of the invention are preferably administered by injection, either subcutaneous or intravenous. The route of administration eventually chosen will depend upon a number of factors and may be ascertained by one skilled in the art.

25

The following examples are offered to more fully illustrate the invention, but are not to be construed as limiting the scope thereof.

30

#### EXAMPLE 1

##### Construction of Hyperglycosylated Epo Analogs

Construction of cDNAs encoding Hyperglycosylated Epo Analogs

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Epo analogs were made by in vitro mutagenesis using several different methods. Analogs N49 and N50 were constructed as described in WO94/09257. Analogs were also constructed by variations of overlap PCR (polymerase chain reaction) methods. The basic procedure included two successive steps. In the first step, two reactions (PCR1 and PCR2) were performed on Epo or Epo analog template DNA using a total of four oligonucleotides: a 5' (forward) primer, a reverse mutagenic primer, a forward mutagenic primer (usually complementary to the reverse mutagenic primer) and a 3' (reverse) primer. The mutagenic primers contained the desired nucleotide changes as well as 6-14 exact match nucleotides on each side of these changes. PCR1 used the 5' (forward) primer and the reverse mutagenic primer. PCR2 used the 3' (reverse) primer and the forward mutagenic primer. The amplified DNA fragments were separated by agarose gel electrophoresis. Small pieces of agarose containing DNA fragments of the correct size were excised from the gel. The DNA fragments from PCR1 and PCR2 were combined together and a third PCR reaction was performed using only the 5' forward and 3' reverse primers. Thus, a full length DNA segment containing the desired mutations was amplified. In several cases, two or three mutations were combined by introducing a new substitution into DNA already containing a change, using the same PCR process. To construct these multiple glycosylation site analogs, single double or triple site analogs (produced as described above) were used as PCR template, and an additional glycosylation site was introduced by site directed mutagenesis with the appropriate primers.

The Epo analogs N51, N52 and N53 were constructed by the overlap PCR (polymerase chain

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reaction) method 1. One additional N-glycosylation site was introduced in each case. N56 added a glycosylation site (N114 T116) to native sequence HuEpo by using pDSR $\alpha$ 2 Epo as PCR template, N51 added an 5 O-linked glycosylation site (Thr125) to N47 Epo by using pDSR $\alpha$ 2 Epo N47 template (Asn30, Thr32, Val87, Asn88, Thr90) and analog N59 added a glycosylation site (Asn53) to analog N47 using pDSR $\alpha$ 2 EpoN47 template.

10 Polymerase chain reactions for method 1 were performed using a protocol adapted from Cheng et. al., (Proc. Natl. Acad. Sci. USA 91, 5695 (1994)). The 3' (reverse) primer contained sequences that introduced a stop codon followed by a Xba I restriction site:

ATCTAGAAGTTGCTCTGGACAGTTCCCT (SEQ ID NO: 2).

15 The 5' forward reaction primer:

GAAGCTTGC GCC ACCATGGGGTGCACGAATG (SEQ ID NO: 3)

had an Hind III restriction site followed by a Kozak sequence upstream of the Epo initiator codon (ATG).

20 The typical PCR reaction mix contained: 4  $\mu$ l each of forward and reverse primers (5 pmol/ $\mu$ l), 1  $\mu$ l template (25 ng), 10  $\mu$ l of 5X LP buffer (100 mM Tricine pH 8.7/25% glycerol/425 mM KOAc), 10  $\mu$ l dNTP stock (1 mM each of dATP, dTTP, dCTP, dGTP), 0.8  $\mu$ l rtTh polymerase (Perkin Elmer; 2.5 U/ $\mu$ l), and 2  $\mu$ l Vent polymerase (NEB; 0.01 U/ $\mu$ l after 1:100 fresh dilution in 1X LP buffer). H<sub>2</sub>O was added to bring the final volume to 50  $\mu$ l. All the components were added together in the order shown and the PCR was started when the temperature during the first cycle was above 60°C by 25 adding 1  $\mu$ l of 50 mM MgOAc. Typical reaction 30

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conditions were: 2 cycles of 94°C, 10 sec/50°C, 1 min./68°C, 5 min. followed by 25 cycles of 94°C, 10 sec/55°C, 1 min./ 68°C, 5 min. The amplified fragments were separated by agarose gel electrophoresis and the 5 correct sized DNA fragment was purified using a Geneclean™ kit and procedures supplied by the manufacturer (Bio 101, Inc.). The purified DNA was digested with Hind III and Xba I, then it was purified again using the Geneclean™ kit. The fragment was then 10 ligated into Hind III and Xba I cut pDSR $\alpha$ 2 vector. Ligated DNA was precipitated with 2 volumes of ethanol in 0.3M NaOAc pH 5.2 in the presence of carrier tRNA and transformed into *E. coli*. Epo analogs were screened by restriction digest on mini DNA preps. Plasmids from 15 positive clones were then prepared and the insert was sequenced to confirm the presence of the desired mutations and to ensure that no additional amino acid changes were introduced.

Analogs N54 to N61 were constructed using 20 overlap PCR strategy method 2. The 3' (reverse) primer contained sequences that introduced a stop codon followed by a XbaI restriction site:

GATCCTCTAGAGTTGCTCTGGACAG (SEQ ID NO: 4).

The 5' forward reaction primer:

25 CAACAAGCTTGCAGCCATGGGG (SEQ ID NO: 5) had a HindIII restriction site followed by a Kozak sequence upstream of the Epo initiator codon (ATG). A high fidelity PCR strategy was performed using Perkin Elmer Ultma DNA Polymerase and accompanying reagents; 30 10  $\mu$ l 10X PCR buffer, 3  $\mu$ l 1 mM dNTPs, 5 pmol of each primer, and water in a final volume of 100  $\mu$ l. 0.5 units of Ultma polymerase was added after the PCR

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mixture reached 94°C. PCR reactions were then carried out for 5 cycles at 94°C for 30 seconds, 50°C for 30 seconds, and 72°C for 90 seconds. A subsequent 25 cycles were performed at 94°C for 30 seconds, and 72°C 5 for 90 seconds. Product bands of the correct sizes were excised from an agarose gel following electrophoresis.

The resulting PCR products for each analog were cleaned using the Qiagen gel extraction kit. The 10 purified DNA was digested in a 100 µl restriction digest with HindIII and XbaI restriction enzymes (Boehringer Mannheim) at 37° C for 1 hour. The digests were again gel purified and the digested fragment was then ligated into HindIII and XbaI digested pDSR $\alpha$ 2 15 vector.

Ligated DNA was precipitated with 2 volumes of ethanol in 0.3M NaOAc pH 5.2 in the presence of carrier tRNA and transformed into E. coli. Epo hyperglycosylated analogs were initially screened by 20 colony PCR to identify clones containing the correctly sized and type of DNA insert. With this procedure, cells containing plasmids were placed into PCR tubes in the presence of Epo forward and reverse primers. The mixture was then subjected to PCR using the reaction 25 conditions described above. Plasmids from positive clones were then prepared and the Epo analog insert was sequenced to confirm the presence of the desired mutations and to ensure that no additional amino acid changes were introduced.

TABLE 1  
ERYTHROPOIETIN ANALOGS HAVING SITES  
FOR N-LINKED CARBOHYDRATE CHAINS

Analog	Amino Acid Substitution	Sequence Changes
N49	Lys, Met → Asn52, Thr54	AAG, ATG → AAT, ACC
N50	Arg, Glu → Asn53, Thr55	AGG, GAG → AAT, ACG
N51	Ala, His, Pro, Trp, Pro, Ala → N30, T32, V87, N88, T90 T125	GCT, CAC, CCG, TGG, CCC, GCC → AAT, ACG, GTG, AAT, ACC, ACC
N52	Ala, Lys → Asn114, Thr116 →	GCC, AAG → AAC, ACG
N53	Ala, His, Arg, Glu Pro, Trp, Pro → N30, T32, N53, T55, V87, N88, T90	GCT, CAC, AGG, GAG, CCG, TGG, CCC → AAT, ACG, AAT, ACG, GTG, AAT, ACC
N54	Glu, Gly → Asn55, Thr57	GAG, GGG → AAT, ACT
N55	Gln, Pro, Trp → Asn86, Val87, Thr88	CAG, CCG, TGG → ACC, GTG, ACG
N56	Pro, Trp, Pro → Ala87, Asn88, Thr90	CCG, TGG, CCC → GCG, AAT, ACC
N57	Pro, Trp, Pro → Val87, Asn88, Ser90	CCG, TGG, CCC → GTG, AAT, ACG
N58	Pro, Trp, Glu, Pro → Val87, Asn88, Gly89, Thr90	CCG, TGG, GAG, CCC → GTG, AAT, GGG, ACC
N59	Ala, His, Arg, Glu → Asn30, Thr32, Asn53, Asn55	GCT, CAC AGG GAG → AAT, ACG, AAT, ACG
N60	Ala, His, Ala, Lys → Asn30, Thr32, Asn114, Thr116	GCT, CAC, GCC, AAG → AAT, ACG, AAC, ACG
N61	A, H, R, E, P, W, P, A, K → I0, T32, N53, T55, V87, N88, T90, N114, T115	GCT, CAC, ACG, GAG, CCG, TGG, CCC, GCC, AAG → AAT, ACG, AAT, A CG, GTG, AAT, ACC, AAC, ACG

Analysis of carbohydrate addition

The constructs for the hyperglycosylated Epo  
5 analogs which were inserted into the expression vector  
pDSR $\alpha$ 2 were transfected into COS cells. Supernatants  
from the transfected COS cells were analyzed by western  
blot to determine whether the expressed and secreted  
Epo analog contained additional carbohydrate. Samples  
10 were loaded directly into wells of SDS-PAGE gels then  
analyzed by immunoblot using the monoclonal antibody,  
9G8A (Elliott et al (1996) Blood 87:p2714). Mobilities  
of analog samples were compared to that of samples

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containing rHuEpo. Figure 1 shows decreased mobility of analogs N53 and N61 compared to analogs N4 (four carbohydrate chains) and N47 (five carbohydrate chains). The mobility is consistent with the presence 5 of six carbohydrate chains for analog N53 and seven carbohydrate chains for analog N61. Data for all hyperglycosylated analogs are shown in Table 2.

In vitro bioassays

10 Media conditioned by COS or CHO cells expressing rHuEpo or analogs were assayed for stimulation of  $^{3}\text{H}$ -thymidine uptake by UT7-Epo cells (Komatsu et al., Blood 82, 456). UT7-Epo cells are responsive to Epo and express human Epo receptors on 15 their cell surface. UT7-Epo cells were grown in Growth medium(1X Iscove's Modified Dulbecco's Medium with L-glutamine, 25 mM HEPES buffer, and 3024 mg/L sodium bicarbonate, but without either alpha-thioglycerol or beta-mercaptoethanol (GIBCO)/ 10% v/v Fetal Bovine 20 Serum/ 1% v/v L-glutamine-Penicillin-Streptomycin solution (Irvine Scientific)/ 1 Unit/mL rHuEpo ) to approximately  $3 \times 10^5$  cells/mL. Cells were collected by centrifugation (approx. 500xG) washed twice with phosphate buffered saline and resuspended at  $5 \times 10^4$  25 cells/mL in Assay medium (1x RPMI Medium 1640 without L-glutamine (Gibco)/1% L-glutamine/4% fetal bovine serum). Test samples or Epo standard (rHuEpo), 100  $\mu\text{L}$  diluted in assay medium at least 5-fold, were added to wells in a 96 well microtiter plate. 50 $\mu\text{L}$  of suspended 30 cells were then added (5000 cells/well) and plates were incubated in a humidified incubator at 37°C and 5% CO<sub>2</sub>. After 72 hours, 50  $\mu\text{L}$  methyl- $^{3}\text{H}$ -Thymidine (1 mCi/mL; 20 Ci/mMole) diluted 1:100 in assay medium was added. Cells were incubated for an additional 4 hours at 37°C

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and 5% CO<sub>2</sub>. Labeled cells were harvested onto glass fiber filtermats, washed with deionized water followed by 2-propanol, dried and counted. Activity was determined by comparing the response determined for 5 each analog to that of the rHuEpo standard. The specific biological activity was then determined by dividing in vitro activity by the concentration of each analog as determined by immunoassay (Elliott et al (1996) Blood 87:p2714). The results are shown in Table 10 2.

TABLE 2

ANALOG	Number of N-linked Carbohydrate Chains	In Vitro Activity **
rHuEpo	3	+++
N49	4	+++
N50	4	+++
N51	5*	+++
N52	3 - 4	+++
N53	6	++
N54	4	NT
N55	4	+++
N56	4	+++
N57	3 - 4	+++
N58	4	+++
N59	5	++
N60	4 - 5	+++
N61	6 - 7	NT

15 \* contains 1-2 O-linked chains  
 \*\* In vitro activity is relative to rHuEpo activity  
 +++ activity equivalent to rHuEpo  
 ++ activity is 25-75% of rHuEpo  
 NT Not Tested

20 Epo analogs N62-N69 were made by overlap PCR (polymerase chain reaction) methods. The basic procedure included two successive steps. In the first step, two reactions (PCR1 and PCR2) were performed on Epo or Epo analog template DNA using a total of four 25 oligonucleotides: a 5' (forward) primer, a reverse

mutagenic primer, a forward mutagenic primer complementary to the reverse mutagenic primer and a 3' (reverse) primer. The mutagenic primers contained the desired nucleotide changes as well as 6-14 exact match 5 nucleotides on each side of these changes. PCR1 used the 5' (forward) primer and the reverse mutagenic primer. PCR2 used the 3' (reverse) primer and the forward mutagenic primer. The amplified DNA fragments were separated by agarose gel electrophoresis and DNA 10 fragments of the correct size were excised and eluted from the gel. The DNA fragments from PCR1 and PCR2 were combined together and a third PCR reaction was performed using only the 5' forward and 3' reverse primers. For some analogs, three PCR reactions were 15 required to generate the desired sequence. These were carried out as above, with a second pair of mutagenic primers being used to generate the third product. Again, the amplified DNA fragments were gel purified and combined in a final reaction containing only the 5' 20 forward and 3' reverse primers. In each case, a full length DNA segment containing the desired mutations was amplified.

N62 added two glycosylation sites (N30 T32 N55 T57) to native sequence HuEpo by using pDSR $\alpha$ 2 Epo 25 N4 (N30 T32) as PCR template. N63 added three glycosylation sites (N30 T32 N55 T57 V87 N88 T90) to native sequence HuEpo by using pDSR $\alpha$ 2 Epo N4 (N30 T32) and pDSR $\alpha$ 2 Epo N47 (N30 T32 N55 T57) as PCR templates. N64 added three glycosylation sites (N30 T32 N55 T57 30 N114 T116) to native sequence HuEpo by using pDSR $\alpha$ 2 Epo N4 (N30 T32) and pDSR $\alpha$ 2 Epo N60 as PCR templates. N65 added three glycosylation sites (N30 T32 V87 N88 T90 N114 T116) to native sequence HuEpo by using pDSR $\alpha$ 2 Epo

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N4 (N30 T32) and pDSR $\alpha$ 2 Epo N60 as PCR templates. N66 added four glycosylation sites (N30 T32 N55 T57 V87 N88 T90 N114 T116) to native sequence HuEpo by using pDSR $\alpha$ 2 Epo N4 (N30 T32) and pDSR $\alpha$ 2 Epo N60 as PCR templates.  
5 N67 added an O-linked glycosylation site (P124 T125 T 126) to N64 Epo. N68 added an O-linked glycosylation site (P124 T125 T 126) to N65 Epo. N69 added an O-linked glycosylation site (P124 T125 T 126) to N66 Epo.

For each analog, the same outside primers  
10 were used. The 3' (reverse) primer contained sequences that introduced a stop codon followed by a Sal I restriction site:

AGGTGGACAGTCGACATTATCTGTCCCCGTGTC (SEQ ID NO:11).

The 5' forward reaction primer:

15 AACAAAGCTTCTAGACCACCATGGGGGTG (SEQ ID NO: 12)  
had a Hind III restriction site followed by a Kozak sequence upstream of the Epo initiator codon (ATG) .

Mutagenic primers were as follows:

N30 T32 mutagenic forward primer

20 ACG ACG GGC TGT AAT GAA ACG TGC AGC TTG (SEQ ID NO: 13)

N30 T32 mutagenic reverse primer

CAA GCT GCA CGT TTC ATT ACA GCC CGT CGT G (SEQ ID NO: 14)

N55 T57 mutagenic forward primer

25 GCC TGG AAG AGG ATG AAT GTC ACGCAG CAG GCC GTA GAA (SEQ ID NO: 15)

N55 T57 mutagenic reverse primer

TTC TAC GGC CTG CTG CGT GAC ATT CAT CCT CTT CCA GGC A (SEQ ID NO: 16)

30 V87 N88 T90 mutagenic forward primer

- 40 -

TCT TCC CAG GTG AAT GAG ACC CTG CAG CTG (SEQ ID NO: 17)

V87 N88 T90 mutagenic reverse primer

CAG CTG CAG GGT CTC ATT CAC CTG GGA AGA GTT G (SEQ ID NO: 18)

5 P124 T125 T126 mutagenic forward primer

CCA GAT CCG ACC ACA GCT GCT CCA (SEQ ID NO: 19)

P124 T125 T126 mutagenic reverse primer

TGG AGC AGC TGT GGT CGG ATC TGG A (SEQ ID NO: 20)

10 The N114 T116 changes were introduced using a template containing the appropriate mutations, so no mutagenic primers were required to generate this site.

The typical PCR1 reaction mix contained: 2.5  $\mu$ l each of forward and mutagenic reverse primers (10 pmol/ $\mu$ l), 1  $\mu$ l template (25 ng), 10  $\mu$ l of 10X Taq Extend buffer (Stratagene), 2  $\mu$ l dNTP stock (10 mM each of dATP, dTTP, dCTP, dGTP), 0.5  $\mu$ l Taq polymerase (BMB), and 0.5  $\mu$ l Taq Extend (Stratagene). H<sub>2</sub>O was added to bring the final volume to 100  $\mu$ l. Typical reaction conditions were: 1 cycle of 94°C, 5 min./55°C, 1 min./ 68°C, 1 min. followed by 25 cycles of 94°C, 1 min./55°C, 1 min./ 68°C, 1 min. The typical PCR2 reaction was identical to that described for PCR 1, except that the reverse and mutagenic forward primers were used. Where a third initial reaction was required, the reaction contained mutagenic forward and mutagenic reverse primers. The amplified fragments were separated by agarose gel electrophoresis and the correct sized DNA fragment was purified using a Gel Extraction kit and procedures supplied by the manufacturer (Qiagen). 30 Complementary fragments were then combined in a third

- 41 -

PCR reaction using only the outside forward and reverse primers. The amplified fragments were separated by agarose gel electrophoresis and purified from the gel as described above. The purified DNA was digested with 5 Hind III and Sal I, then it was again gel purified. The fragment was then ligated into Hind III and Sal I cut pDSR $\alpha$ 19 vector. Ligated DNA transformed by electroporation into *E. coli*. Epo hyperglycosylated analogs were initially screened by colony PCR to 10 identify clones containing the correctly sized DNA insert. Plasmid DNA from selected clones was then prepared and the insert was sequenced to confirm the presence of the desired mutations and to ensure that no additional amino acid changes were introduced.

15

TABLE 3  
ERYTHROPOIETIN ANALOGS HAVING SITES  
FOR N-LINKED CARBOHYDRATE CHAINS

Analog	Amino Acid Substitution	Sequence Changes
N62	Ala, His, Glu, Gly $\rightarrow$ Asn30, Thr32, Asn55 Thr57	GCT, CAC, GAG, GGG $\rightarrow$ AAT, ACG, AAT, ACT
N63	A, H, E, G, P, W, P $\rightarrow$ N30, T32, N55, T57, V87, N88, T90	GCT, CAC, GAG, GGG, CCG, TGG, CCC $\rightarrow$ AAT, ACG, AAT, ACT GTG, AAT, ACC
N64	A, H, E, G, A, K $\rightarrow$ N30, T32, N55, T57, N114 T116	GCT, CAC, GAG, GGG, GCC, AAG $\rightarrow$ AAT, ACG, AAT, ACT, AAC, ACG
N65	A, H, P, W, P, A, K $\rightarrow$ N30, T32, V87, N88, T90, N114, T116	GCT, CAC, CCG, TGG, CCC, GCC, AAG $\rightarrow$ AAT, ACG, GTG, AAT, ACC, AAC, ACG
N66	A, H, E, G, P, W, P, A, K $\rightarrow$ N30, T32, N55, T57, V87, N88, T90, N114, T116	GCT, CAC, GAG, GGG, CCG, TGG, CCC, GCC, AAG $\rightarrow$ AAT, ACG, AAT, ACT GTG, AAT, ACC, AAC, ACG
N67	A, H, E, G, P, W, P, A, A, S $\rightarrow$ N30, T32, N55, T57, V87, N88, T90, P124, T125, T126	GCT, CAC, GAG, GGG, CCG, TGG, CCC, GCC, GCC, TCA $\rightarrow$ AAT, ACG, AAT, ACT GTG, AAT, ACC, CCG ACC ACA
N68	A, H, E, G, A, K, A, A, S $\rightarrow$ N30, T32, N55, T57, N114 T116, P124, T125, T126	GCT, CAC, GAG, GGG, GCC, AAG, GCG, GCC, TCA $\rightarrow$ AAT, ACG, AAT, ACT, AAC, ACG, CCG, ACC, ACA

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N69	A, H, E, G, P, W, P, A, K, A, A, S→ N30, T32, N55, T57, V87, N88, T90, N114, T116, P124, T125, T126	GCT, CAC, GAG, GGG, CCG, TGG, CCC, GCC, AAG, GCG, GCC, TCA→ AAT, ACG, AAT, ACT, GTG, AAT, ACC, AAC, ACG, CCG, ACC, ACA
N70	Ala, His, Pro, Trp, Pro → N30, T32, V87, N88, T90, IgG1 fusion	GCT, CAC, CCG, TGG, CCC→AAT, ACG, GTG, AAT, ACC

Construction of cDNA encoding Hyperglycosylated Epo Analog Fusion Polypeptide

5                   Epo analog N70 was also made by overlap PCR. Plasmid DSRα2 containing the cDNA sequence encoding analog N47 (N30 T32 V87 N88 T90) and plasmid pAMG21 (ATCC accession no. 98113) containing cDNA encoding an Fc region were used as templates for the polymerase 10 chain reactions. The Fc portion of human immunoglobulin IgG1 heavy chain from residue 104 of the hinge domain(Asp-104) to the carboxyl terminus (Ellison et al., *supra*, see also Figure 10 starting at aspartic acid residue at position 6), was generated by PCR 15 amplification of a human spleen cDNA library (Clontech). Overlapping PCR products were generated in two reactions using the following oligonucleotide primers

5' forward reaction primer 2343-85 (Epo specific) :  
20 AAC AAG CTT CTA GAC CAC CAT GGG GGT G (SEQ ID NO: 21)  
3' reverse reaction primer 2343-87 (homology to both Epo and Fc) :  
AGG TGG ACA TGT GTG AGT TTT GTC TCT GTC CCC TCT CCT GCA  
GGC CTC C (SEQ ID NO: 22)  
25 5' forward reaction primer 2343-86 (homology to both Epo and Fc) :

GAG GCC TGC AGG ACA GGG GAC AGA GAC AAA ACT CAC ACA TGT  
CCA CCT (SEQ ID NO: 23)

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3' reverse reaction primer 2343-88 (specific to Fc):  
TGG ACA GTC GAC ATT ATT TAC CCG GAG ACA GGG AGA GGC TCT  
TCT GC (SEQ ID NO: 24)

5 PCR1 contained 2.5 µl each of forward (2343-85) and  
reverse (2343-87) primers (10 pmol/µl), while PCR2  
contained 2.5 µl each of forward (2343-86) and reverse  
(2343-88) primers (10 pmol/µl). Conditions were as  
described above. The resulting amplified products  
10 contained a region of overlap (48 nucleotides) encoding  
the last 8 amino acids of Epo and the first 8 amino  
acids of Fc. The complementary fragments were gel  
purified and combined in a third PCR reaction using  
only the outside forward and reverse primers. The  
15 amplified fragment was separated by agarose gel  
electrophoresis and purified from the gel as described  
above. The purified DNA was digested with Hind III and  
Sal I, then it was again gel purified. The fragment  
was then ligated into Hind III and Sal I cut pDSR $\alpha$ 19  
20 vector. Ligated DNA transformed by electroporation into  
*E. coli*. Transformants were initially screened by  
colony PCR to identify clones containing the correctly  
sized DNA insert. Plasmid DNA from selected clones was  
then prepared and the insert was sequenced to confirm  
25 the sequence of the fusion protein and to ensure that  
no additional amino acid changes were introduced.

Analysis of carbohydrate addition

The constructs for the hyperglycosylated Epo  
30 analogs N62 to N69 and fusion protein (analog N70) were  
inserted into the expression vector pDSR $\alpha$ 19 and  
transfected into CHO cells. Supernatants from the  
transfected CHO cells are analyzed by western blot to

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determine whether the expressed and secreted Epo analogs contained additional carbohydrate using procedures described above for analogs N49 to N62.

5 In vitro bioassays

In vitro assays for analogs N62 to N70 expressed in CHO transfected cells are performed as described above for analogs N49 to N61.

10

EXAMPLE 2

Preparation of Recombinant Human Erythropoietin and Hyperglycosylated Erythropoietin Analogs

15 Recombinant human erythropoietin (rHuEpo) used for the experiments described herein was expressed by Chinese hamster ovary (CHO) cells transfected with a recombinant plasmid carrying the human erythropoietin gene. The recombinant product was recovered from the conditioned medium and purified essentially as 20 described by Lai et al. *supra*. The resulting rHuEpo preparation has predominantly isoforms of 9 to 14 sialic acids as determined by isoelectric focusing.

25 Recombinant hyperglycosylated erythropoietin analogs were expressed in CHO cells transfected with a recombinant plasmid carrying the Epo analog gene as described WO91/05867 and WO94/09257 hereby incorporated by reference. The hyperglycosylated analogs were purified from culture supernatants as described below.

30 Concentration and diafiltration of conditioned media

Conditioned medium (serum free) from three successive harvests (5-8 days each) of the transfected CHO cell line was collected, filtered through a 0.45  $\mu$ m filter, concentrated about thirty fold, and diafiltered

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into 10 mM Tris, 20  $\mu$ M CuSO<sub>4</sub>, pH 7.0 using a tangential-flow ultrafiltration system (Millipore) with a 10,000 molecular weight cutoff membrane. The dialfiltered media (DFM) was filtered (0.45  $\mu$ m) a 5 second time and stored at -20°C until used for purification.

#### Purification

All procedures were carried out at 2 to 8°C.

10

#### Anion - Exchange Chromatography (10)

The clarified DFM was applied to a Q-Sepharose Fast Flow column (Pharmacia, 6 cm x 18 cm) equilibrated in 10 mM bis Tris propane (BTP), pH 7.0 15 and washed with two column volumes of 10 mM BTP to elute all non-binding species. The following gradients were run depending upon whether the hyperglycosylated analog had four, five or six N-linked carbohydrate chains. All buffers used at this stage contain 1 mM glycine, 20  $\mu$ M CuSO<sub>4</sub>, 6 M urea, 5  $\mu$ g/mL leupeptin and 20 1  $\mu$ g/mL pepstatin. For analogs with four N-linked carbohydrate chains, the gradient was 10 mM acetic acid, 0.1 mM NaCl to 500 mM acetic acid, 5 mM NaCl over 49 column volumes with a two column volume hold at high 25 salt conditions. For analogs with five N-linked carbohydrate chains, the gradient was 0.7 M acetic acid, 7 mM NaCl to 1.0 M acetic acid, 12 mM NaCl over 30 column volumes with a two column volume hold at high salt conditions. For analogs with six N-linked 30 carbohydrate chains, the gradient was 1.0 M acetic acid, 10 mM NaCl to 1.5 M acetic acid, 20 mM NaCl over 50 column volumes with a two column volume hold at high salt conditions. Following the gradient, the column was washed with two column volumes of 10 mM BTP, pH 7.0 and

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the high isoform fraction was eluted with 0.6 M NaCl, 100 mM BTP, pH 7.0.

5

Reversed Phase Chromatography (C4)

The high salt strip from the Q-Sepharose column (1Q) was applied to a Vydac C4 reversed phase column (30  $\mu$  particles, 4 cm x 18 cm) equilibrated in 10 20% ethanol, 10 mM BTP, pH 7.0 and eluted from the column with a thirty column volume gradient to 94% ethanol buffered in 10 mM BTP, pH 7.0. The pooled product peak, eluting in approximately 60% ethanol, was diluted with four volumes of 10 mM BTP, pH 7.0 to 15 minimize possibility of aggregation in the presence of ethanol.

Anion - Exchange Chromatography (20)

The diluted eluate from the reversed phase column was applied to a second Q-Sepharose Fast Flow (Pharmacia, 3 cm x 9 cm) column equilibrated with 10 mM BTP, pH 7.0. The column was washed with equilibration buffer, and the hyperglycosylated Epo analog was eluted with 0.6 M sodium chloride, 20 mM sodium citrate, pH 25 6.0.

The purified protein was exchanged into 20 mM NaPO<sub>4</sub>, pH 6.0, 140 mM NaCl via centricon (10,000 molecular weight cutoff), followed by passage through a 0.2  $\mu$ m filter and storage at 2-8°C.

30

EXAMPLE 3

In Vivo Bioactivity of rHuEpo and rHuEpo Analogs  
Containing four, five and six N-linked  
35 Carbohydrate Chains

The in vivo activity of Epo analogs containing four, five and six N-linked carbohydrate chains was compared with that of rHuEpo in the 5 exhypoxic polycythemic mouse bioassay. This assay quantifies the incorporation of <sup>59</sup>Fe into newly synthesized red blood cells as a measure of the increase in erythropoiesis in mice in response to an exogenously-administered test sample. The assay, 10 performed as described below, is a modification of the method of Cotes and Bangham (Nature 191, 1065 (1961)).

In this assay, female BDF<sub>1</sub> mice are first preconditioned by exposure to low oxygen conditions in a hypobaric chamber (0.4 -0.5 atm.) for approximately 15 18 hours a day for 14 days. To compensate for the low oxygen conditions the mice respond by stimulating erythropoiesis to increase the number of red blood cells and thereby the relative oxygen-carrying capacity. After completion of the final hypobaric 20 exposure, the mice are allowed to remain at ambient pressure for approximately 72 hours prior to the administration of test samples by intraperitoneal injection. At ambient pressure the mice are relatively polycythemic and respond by decreasing endogenous 25 erythropoietin production and the rate of erythropoiesis. Five days after sample administration, 0.2 -0.3 µCi of <sup>59</sup>FeCl<sub>3</sub> in 0.2 mL is injected intravenously into the tail vein. Forty-eight hours later, the animals are sacrificed and the increase in 30 erythropoiesis produced by the test samples is determined by measuring the amount of <sup>59</sup>Fe incorporated in a 0.5 mL sample of whole blood.

An example of the results obtained when rHuEpo and five different Epo analogs, containing four, 35 five or six N-linked carbohydrate chains were tested in

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this assay is shown in Figure 3. Each sample was assayed at six or seven different dilutions within an appropriate concentration range. All samples were diluted in phosphate-buffered saline containing 0.5% 5 bovine serum albumin, and 0.4 mL of each dilution was administered to five preconditioned mice. Forty-eight hours after the administration of <sup>59</sup>Fe, the amount incorporated in 0.5 mL of blood was measured by gamma counting. The results for each of the samples are 10 plotted as the percent <sup>59</sup>Fe incorporated versus the log of the administered dose.

As shown in Figure 3, all five of the hyperglycosylated Epo analogs tested in this assay were more potent than rHuEpo. In addition, the potency of 15 each analog was directly dependent on the number of N-linked carbohydrate chains, with those analogs having an increased number of carbohydrate chains having the greater activity. Thus, analog N53, which contains six N-linked carbohydrate chains, was the most potent 20 analog. Analog N47, which contains five N-linked carbohydrate chains, was in turn, more potent than those analogs containing four N-linked chains. The 25 potencies of the three analogs containing four N-linked carbohydrate chains (N4, N18 and N50) were approximately equal to each other and greater than that of rHuEpo.

In this experiment, the doses of rHuEpo and 30 analogs containing four, five or six N-linked carbohydrate chains required to produce 40% <sup>59</sup>Fe incorporation were 10,700 ng, 640 ng, 140 ng and 38 ng, respectively. Based on the amount of material required to produce this level of erythropoiesis, Epo analogs containing four, five, or six N-linked carbohydrate chains are 17-fold, 77-fold and 280-fold more potent 35 than rHuEpo.

5

## EXAMPLE 4

IV Pharmacokinetics of rHuEpo and Epo analog  
N47 in Rats and Beagle Dogs

10

Two separate studies in rats and dogs were performed to compare the pharmacokinetic parameters of Epo N47 analog and rHuEpo.

In the rat studies, 1  $\mu$ Ci ( $\sim$ 0.1  $\mu$ g of peptide/kg) of either  $^{125}$ I-Epo N47 analog, or  $^{125}$ I-recombinant human erythropoietin (Amersham) was injected intravenously into a surgically implanted carotid cannula in normal male Sprague-Dawley rats weighing between 314-363 g. At various time points after administration, 0.3mL of blood was collected and serum was prepared by centrifugation. The level of  $^{125}$ I-rHuEpo or  $^{125}$ I Epo N47 analog in 0.1mL of each serum sample was then determined following an overnight 4°C incubation with 90% ethanol. The ethanol-precipitated protein in each serum sample was collected by centrifugation and the radioactivity was counted in a gamma counter. The resulting serum concentration vs time pharmacokinetic curves are shown in Figure 4. Each point represents a group mean of five rats in the N47 analog group and six rats in the rHuEpo group. The pharmacokinetic parameters were determined for each rat using PCNONLIN 4.0 nonlinear regression analysis (Statistical Consultants, 1992) and the

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results for each group were averaged. The results are shown in Table 3.

5

TABLE 3

10 Comparison of IV Pharmacokinetic Parameters of N47 and r-HuEpo in Rats

Sample Test	Half-life		V <sub>d</sub> (mL/kg)	Serum Clearance (mL/kg-hr)
	$\alpha$ (hours)	$\beta$ (hours)		
N47 (n = 5 rats)	0.57±0.49	6.9±0.3	33±5	4.8±1.2
r-HuEpo (n = 6 rats)	0.18±0.03	2.5±0.2	36±6	17.7±3.4

15 a The results are presented as the group average  $\pm$  SD for five rats in the N47 group and six rats in the r-HuEpo group.

In the dog studies, normal Beagle dogs 20 weighing between 7.8-9.5 kg received an intravenous bolus injection of ~29 $\mu$ Ci of either <sup>125</sup>I-rHuEpo or <sup>125</sup>I-N47 (~0.1 $\mu$ g peptide/kg) into the cephalic vein. At various time points through 24 hours post-administration, approximately 1 to 2 mL of blood was 25 collected and serum prepared. The concentration of <sup>125</sup>I-rHuEpo and <sup>125</sup>I-N47 in 0.1mL serum was determined and pharmacokinetic parameters were calculated as described above. The serum concentration vs time pharmacokinetic curves for the dog studies are shown 30 in Figure 5. The time points are the group means of two animals in each group. The pharmacokinetic parameters are summarized in Table 4.

35

TABLE 4

Comparison of IV Pharmacokinetic Parameters of N47

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and r-HuEpo in Dogs

Sample Test	Half-life		V <sub>a</sub> (mL/kg)	Serum Clearance (mL/kg-hr)
	α (hours)	β (hours)		
N47 r-HuEpo	0.34 0.40	25.0 7.2	55.9 60.8	2.4 8.4

5

a The results presented are the average parameters for the two dogs in each group.

10 In both the rat and dog studies, rHuEpo and Epo N47 analog exhibited a biphasic serum clearance. Clearance in rats was about 3.7-fold faster for rHuEpo than for Epo N47 analog and the β-half-life was about 2.8-fold longer for Epo N47 analog than for rHuEpo. The pharmacokinetic 15 parameters in the dog studies were generally consistent with those observed in rat. In dogs, the clearance of rHuEpo was 3.5-fold faster than for Epo N47 analog and the β-half-life was 3.5-fold longer for Epo N47 analog compared with that for rHuEpo.

20

#### EXAMPLE 5

Dose response of hematocrit after administration of rHuEpo and Epo analog N47.

25 Hematocrit Dose Response Studies at three times per week (TIW)

30 The in vivo biological effects of rHuEpo and Epo analog N47 in normal mice were compared after administering a range of doses by either intraperitoneal or intravenous injection three times per week for up to six weeks. Hematocrit determinations were performed twice weekly by retro-orbital bleed.

35 Normal CD1 mice weighing approximately 30g (10-13 mice per group) were injected

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intraperitonally three times per week for a total of six weeks with either rHuEpo (over the dose range of 0.625-10 $\mu$ g peptide/kg/dose), Epo N47 analog (over the dose range of 0.156-1.25 $\mu$ g peptide/kg/dose) or 5 vehicle control. The vehicle control and diluent for the various rHuEpo and Epo N47 analog dosing preparations was phosphate-buffered saline (PBS), containing 0.025% mouse serum albumin. The hematocrits of all mice were determined at baseline 10 and twice weekly thereafter by retro-orbital bleeds. At the conclusion of the experiment, serum from all animals was collected and assayed for antibodies to the injected product by a solution radioimmunoprecipitation assay. Hematocrit data 15 from animals judged to be negative for neutralizing antibodies were used for subsequent analysis.

As shown in Figure 6 both rHuEpo and Epo N47 analog produce a dose-dependent increase in hematocrit in the six week study, although N47 analog promotes a 20 greater increase in hematocrit compared to rHuEpo at a given concentration. In this experiment the Epo N47 analog is about 3 to 4-fold more potent when dosed three times per week by intraperitoneal injection.

Dose response studies of rHuEpo and analog 25 N47 were carried out by intravenous injection three times per week using procedures similar to those for intraperitoneal injection. The results obtained were similar to those for intraperitoneal administration and, in particular, the studies further confirmed that 30 Epo N47 analog had a greater potency than rHuEpo when administered three times per week.

To better compare and quantify the biological 35 activity of rHuEpo and Epo N47 analog in raising the hematocrit of normal mice, results of experiments were also analyzed by relative potency plots. For each

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experiment, the activity of rHuEpo or N47 analog at each dose was determined by summing the increase in hematocrit over the first 38 days of the study by trapezoidal summation to obtain the area under the 5 curve (AUC). This was then plotted versus the log dose in  $\mu$ g peptide/kg/week. Potency difference between compounds administered by the same or different routes of administration or dosing frequencies can be determined by measuring the distance between the 10 relevant log-dose response lines. Figure 7 summarizes the relative potency data for all experiments performed comparing the activity of rHuEpo and Epo N47 analog administered by two different routes (intraperitoneal and intravenous) and at two different dosing schedules.

15 As shown in Figure 7 when administered three times per week, Epo N47 analog has the same potency when injected by either the intravenous or intraperitoneal route and was 3.6-fold more potent than rHuEpo injected intraperitoneally three times weekly.

20

Hematocrit Dose Response Studies at one time per week (OW)

25 Comparisons of rHuEpo and Epo analog N47 at increasing hematocrit in normal mice were undertaken with once weekly dosing by either the intraperitoneal or intravenous routes of administration for six weeks.

Normal CD1 mice weighing approximately 30g (8-10 mice per group) were injected intravenously once weekly for a total of six weeks with varying 30 concentrations of either rHuEpo or Epo N47 analog prepared in PBS containing 0.025% mouse serum albumin, or with vehicle control (PBS with 0.025% mouse serum albumin). The analog dose varied from 6.25-25  $\mu$ g of peptide/kg/dose and the dose of rHuEpo 35 varied from 25-200  $\mu$ g/kg/dose. The hematocrits of

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all mice were determined at baseline and twice weekly thereafter by retro-orbital bleeds. At the conclusion of the experiment, serum from all animals was collected and assayed for antibodies to the 5 injected product by a solution radioimmunoprecipitation. Data from animals judged to be negative for neutralizing antibodies were used for subsequent analysis.

As shown in Figure 8, whereas both rHuEpo and 10 analog N47 can increase the hematocrit of normal mice when dosed once weekly, the dose of rHuEpo required to produce a response was significantly greater than that for analog N47. For instance, in this experiment 25 $\mu$ g peptide/kg/week of N47 increased the hematocrit of mice 15 by 41.2 points in six weeks, whereas the same dose of rHuEpo produced only a 12.5 point hematocrit rise.

Dose response studies of rHuEpo and analog 20 N47 were performed by intraperitoneal injection once weekly using procedures similar to those described above. The results obtained were consistent with the results for intravenous administration and further confirmed the greater potency of analog N47 compared with rHuEpo when administered one time per week.

To quantify the activity difference between 25 rHuEpo and N47 analog when each is dosed once weekly, relative potency plots were generated from all relevant experiments as described above. As shown in Figure 7, when administered one time per week, analog N47 has the same potency when injected by the intravenous and 30 intraperitoneal route. Analog N47 is approximately 14-fold more potent than rHuEpo when each is administered once weekly.

In addition, the log-dose response plots in Figure 6 also illustrate the following: (1) A given 35 dose of analog N47 administered once weekly (QW) is

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approximately as effective as the same total weekly dose of rHuEpo given as three divided doses (TIW); (2) A given dose of rHuEpo administered once weekly (QW) is only approximately 2% as effective as the same total 5 weekly dose of analog N47 given as three divided doses (TIW); (3) Analog N47 is approximately 4-fold more potent in mice when administered TIW compared to QW.

Hematocrit Dose Response Studies at Every Other Week

10 (EOW)

Experiments were also undertaken to assess the ability of analog N47 to increase the hematocrit of mice when injected once every other week. Normal CD-1 mice (10 mice per group) were injected intravenously 15 either once weekly or once every other week for a total of approximately six weeks with varying concentrations of Epo N47 analog prepared in PBS containing 0.025% mouse serum albumin. Analog N47 was administered at either 200, 100 or 25 $\mu$ g/kg/dose every other week or at 20 12.5 $\mu$ g/kg/dose once weekly. The hematocrits of all mice were determined at baseline and twice weekly thereafter by retroorbital bleeds.

As shown in Figure 9, analog N47 can increase the hematocrit of normal mice in a dose-dependent 25 fashion even when administered bi-monthly. As expected when dosed less frequently, a greater amount of N47 analog is required to increase the hematocrit. A dose of 200  $\mu$ g/kg of N47 analog administered every other week increased the hematocrit to approximately the same 30 extent in six weeks as did 12.5  $\mu$ g/kg when dosed weekly.

EXAMPLE 6

IV Pharmacokinetics of Epo N47 analog

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and rHuEpo in Continuous Ambulatory Peritoneal Dialysis (CAPD) patients

In view of the marked increase in serum  
5 half-life of Epo N47 analog compared to rHuEpo in rat  
and beagle dog, it was of interest to determine whether  
an increase could also be observed in humans.

A double-blind, randomized cross-over design  
study of eleven stable CAPD patients (7 males, 4  
10 females, aged 27-75 years) was undertaken. One patient  
group received 100U/kg of rHuEpo (equivalent to 0.5 µg  
of peptide/kg) while a second group of patients  
received 0.5 µg peptide/kg of Epo N47 analog, both  
administered as a single bolus injection intravenous.  
15 Venous blood samples (3mL) were drawn via an indwelling  
cannula and were taken pre-dose and at 5, 10, 15, 30  
minutes and 1, 2, 5, 8, 12, 16, 24, 30, 36, 48, 60, 72  
and 96 hours after the intravenous bolus. After a 28  
day washout period, the first patient group received a  
20 single intravenous dose of Epo analog N47 while the  
second group received a single intravenous dose of  
rHuEpo. Blood samples were taken as in the first cycle  
of treatment. Levels of rHuEpo and Epo N47 analog were  
determined in serum by ELISA after subtraction of  
25 baseline endogenous Epo levels. The pharmacokinetic  
parameters (mean ± SE) estimated after adjustment for  
cross-over design effects are shown in Table 5. The  
serum concentration AUC was calculated using the linear  
trapezoidal summation.  $t_{1/2}$  is defined as:  $\log(2)/K_z$ ,  
30 where  $K_z$  is calculated as the slope of the terminal  
portion of the ln (serum concentration) time curve.  
The clearance (Cl) is defined as: dose/AUC. The volume  
of distribution ( $V_d$ ) is defined as: Cl/K.

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Dose Group	AUC (ng.h/mL)	$t_{1/2z}$ (h)	Cl (mL/h/kg)	Vd (mL/kg)
N47	291.0±7.6	25.3±2.2	1.6±0.3	52.4±2.0
rHuEpo	131.9±8.3	8.5±2.4	4.0±0.3	48.7±2.1

5                   The mean serum half-life for Epo N47 analog (25.3 hr) was three times longer than for rHuEpo (8.5 hr) and the clearance was 2.5-fold faster for rHuEpo than for analog N47.

10

#### EXAMPLE 7

##### A Phase II Dose Finding and Dose Scheduling Study of Epo N47 Analog

15                   Multicenter, randomized, sequential dose-escalation studies are initiated to investigate the optimum dose and dose schedule for analog N47 when administered by subcutaneous or intravenous injection in patients with CRF receiving dialysis.

20                   The dosing schedule is as follows:

Once per week dosing: 0.075, 0.225, 0.45, 0.75, 1.5 and 4.5  $\mu$ g of peptide/kg/dose.

Three times per week dosing: 0.025, 0.075, 0.15, 0.25, 0.5 and 1.5  $\mu$ g of peptide/kg/dose.

25

                  The studies are carried out in two parts: the first part is a dose-escalation study designed to evaluate the dose of analog N47 given either one or three times per week which increases hemoglobin at an optimum rate over four weeks (greater than or equal to 1 g/dL but less than 3 g/dL). The second part of each study is designed to determine the

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doses required (when administered once or three times per week by either the intravenous or subcutaneous routes of administration) to maintain the hematocrit at the therapeutic target.

5           Preliminary results indicate that once weekly dosing with analog N47 can be used to both increase and maintain the hematocrit of anemic CRF patients. Initial results suggest that the preferred doses to initiate therapy on a three times  
10   a week dosing schedule are 0.15 and 0.25  $\mu$ g/peptide/kg/dose, and on a one time per week dosing schedule are 0.45 and 0.75  $\mu$ g/peptide/kg/dose for both routes of administration.

15

\*       \*       \*

While the invention has been described in what is considered to be its preferred embodiments, it is not to be limited to the disclosed  
20   embodiments, but on the contrary, is intended to cover various modifications and equivalents included within the spirit and scope of the appended claims, which scope is to be accorded the broadest  
25   interpretation so as to encompass all such modifications and equivalents.

## WHAT IS CLAIMED IS:

1. A method of raising and maintaining hematocrit in a mammal comprising administering a 5 therapeutically effective amount of a hyperglycosylated analog of erythropoietin in a pharmaceutical composition, wherein the analog is administered less frequently than an equivalent molar amount of recombinant human erythropoietin to obtain a comparable 10 target hematocrit.

2. The method of Claim 1 wherein the amount of hyperglycosylated analog of erythropoietin is administered about two times per week.

15 3. The method of Claim 1 wherein the amount of hyperglycosylated analog of erythropoietin is administered about one time per week.

20 4. The method of Claim 1 wherein the amount of hyperglycosylated analog of erythropoietin is administered about one time every other week.

25 5. The method of Claim 1 wherein the amount of hyperglycosylated analog of erythropoietin is administered about one time per month.

30 6. The method of Claim 3 wherein the amount of hyperglycosylated analog of erythropoietin that is administered is about 0.075 to 4.5  $\mu$ g per erythropoietin peptide per kg per dose.

35 7. A method of raising and maintaining hematocrit in a mammal comprising administering a therapeutically effective amount of a hyperglycosylated

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analog of erythropoietin in a pharmaceutical composition, wherein the analog is administered at a lower molar amount than recombinant human erythropoietin to obtain a comparable target  
5 hematocrit.

8. The method of Claim 7 whereas the amount of hyperglycosylated analog of erythropoietin that is administered is about 0.025 to 1.5  $\mu$ g erythropoietin  
10 peptide per kg per dose three times per week.

9. The method of Claims 1 or 7 wherein the hyperglycosylated analog of erythropoietin comprises at least one additional glycosylation site compared to  
15 human erythropoietin wherein a carbohydrate chain is added to the site.

10. The method of Claims 1 or 7 wherein the target hematocrit is at least about 30%.

20 11. The analog of Claim 9 wherein the carbohydrate chain is an N-linked carbohydrate chain at one or more of positions 30, 51, 57, 69, 88, 89, 136 and 138 of the sequence of human erythropoietin.

25 12. The analog of Claim 9 which has additional N-linked carbohydrate chains at position 30 and 88 of the sequence of human erythropoietin.

30 13. The analog of Claim 12 which is Asn<sup>30</sup>Thr<sup>32</sup>Val<sup>87</sup> Asn<sup>88</sup>Thr<sup>90</sup> Epo.

14. The method of Claims 1 or 7 wherein the pharmaceutical composition comprises a pharmaceutically

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acceptable diluent, carrier, solubilizer, emulsifier, preservative, and/or adjuvant.

15. The composition of Claim 14 wherein the  
5 diluent is a buffer solution of sodium citrate or  
sodium phosphate.

16. The composition of Claim 14 wherein the  
carrier is human serum albumin.

10

17. The composition of Claim 14 wherein the  
preservative is benzyl alcohol.

18. The method of Claims 1 or 7 wherein the  
15 mammal suffers from anemia associated with a decline or  
loss of kidney function.

19. The method of Claims 1 or 7 wherein the  
mammal suffers from anemia associated with  
20 myelosuppressive therapy.

20. The method of Claim 19 wherein the  
myelosuppressive therapy comprises chemotherapeutic or  
anti-viral drugs.

25

21. The method of Claims 1 or 7 wherein the  
mammal suffers from anemia associated with excessive  
blood loss.

30

22. The method of Claims 1 or 7 further  
comprising administering a therapeutically effective  
amount of iron.

35 23. An analog of human erythropoietin  
comprising at least one additional glycosylation site

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at any of positions 52, 53, 55, 86 and 114 of the sequence of human erythropoietin, wherein an N-linked carbohydrate chain is added to the site.

5 24. The analog of Claim 22 comprising at least two additional glycosylation sites wherein a carbohydrate chain is attached to each of the sites.

10 25. The analog of Claim 22 comprising at least three additional glycosylation sites wherein a carbohydrate chain is attached to each of the sites.

15 26. The analog of Claim 22 comprising at least four additional glycosylation sites wherein a carbohydrate chain is attached to each of the sites.

27. An analog of human erythropoietin selected from the group consisting of:

20 Asn<sup>52</sup> Thr<sup>54</sup> Epo;  
Asn<sup>53</sup> Thr<sup>55</sup> Epo;  
Asn<sup>30</sup> Thr<sup>32</sup> Val<sup>87</sup> Asn<sup>88</sup> Thr<sup>90</sup> Thr<sup>125</sup> Epo;  
Asn<sup>114</sup> Thr<sup>116</sup> Epo;  
Asn<sup>30</sup> Thr<sup>32</sup> Asn<sup>53</sup> Thr<sup>55</sup> Val<sup>87</sup> Asn<sup>88</sup> Thr<sup>90</sup> Epo;  
Asn<sup>55</sup> Thr<sup>57</sup> Epo;  
25 Asn<sup>86</sup> Val<sup>87</sup> Thr<sup>88</sup> Epo;  
Ala<sup>87</sup> Asn<sup>88</sup> Thr<sup>90</sup> Epo;  
Val<sup>87</sup> Asn<sup>88</sup> Ser<sup>90</sup> Epo;  
Val<sup>87</sup> Asn<sup>88</sup> Gly<sup>89</sup> Thr<sup>90</sup> Epo;  
Asn<sup>30</sup> Thr<sup>32</sup> Asn<sup>53</sup> Thr<sup>55</sup> Epo;  
30 Asn<sup>30</sup> Thr<sup>32</sup> Asn<sup>114</sup> Thr<sup>116</sup> Epo; and  
Asn<sup>30</sup> Thr<sup>32</sup> Asn<sup>53</sup> Thr<sup>55</sup> Val<sup>87</sup> Asn<sup>88</sup> Thr<sup>90</sup> Asn<sup>114</sup> Thr<sup>116</sup> Epo.

28. The analog of Claims 23-27 which is the  
35 product of expression of an exogenous DNA sequence.

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29. A DNA sequence encoding the analog of  
Claims 23-27.

5 30. A eucaryotic host cell transfected with  
the DNA sequence of Claim 29 in a manner allowing the  
host cell to express the analog.

10 31. A composition comprising a  
therapeutically effective amount of the analog of  
Claims 23-27 together with a pharmaceutically  
acceptable diluent, adjuvant or carrier.

15 32. A method of raising and maintaining  
hematocrit in a mammal comprising administering a  
therapeutically effective amount of the analog of  
Claims 23-27 in a pharmaceutical composition.

20 33. The method of Claim 32 wherein the  
analog is administered less frequently than an  
equivalent molar amount of recombinant human  
erythropoietin to obtain a comparable target  
hematocrit.

25 34. The method of Claim 33 wherein the  
amount of analog is administered about one time per  
week, about one time every other week, or about one  
time per month.

30 35. The method of Claim 32 wherein the  
analog is administered at a lower molar amount than  
recombinant human erythropoietin to obtain a comparable  
target hematocrit.

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36. The method of Claim 34 wherein the amount of analog that is administered is about 0.025 to 1.5  $\mu$ g erythropoietin peptide per kg per dose three times per week.

5

37. The method of Claim 34 wherein the amount of analog that is administered is less than about 0.025  $\mu$ g erythropoietin peptide per kg per dose three times per week.

10

38. A fusion protein comprising a hyperglycosylated analog of erythropoietin and an immunoglobulin heavy chain constant region.

15

39. The fusion protein of claim 38 wherein the immunoglobulin heavy chain constant region is an Fc region.

20

40. The fusion protein of Claim 39 wherein the Fc region is from human IgG or derived from human IgG.

25

41. The fusion protein of Claim 40 wherein the IgG is IgG1, IgG2, IgG3 or IgG4, or a combination thereof.

30

42. The fusion protein of Claim 38 comprising at least one additional glycosylation site at any of positions 30, 51, 52, 53, 55, 57, 69, 86, 88, 89, 114, 136 and 138 of the sequence of human erythropoietin, wherein an N-linked carbohydrate chain is added at the site.

35

43. The fusion protein of Claim 38 which is Asn<sup>30</sup>Thr<sup>32</sup>Val<sup>87</sup>Asn<sup>88</sup>Thr<sup>90</sup> Epo fused to an Fc region.

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44. The fusion protein of Claim 43 consisting of the mature amino acid sequence as set forth in Figure 11 (SEQ ID NO: 26) lacking the signal sequence.

## FIG.1

Met Gly Val His Glu Cys Pro Ala Trp Leu Trp Leu Leu Leu Ser Leu  
                   -25                  -20                  -15

Leu Ser Leu Pro Leu Gly Leu Pro Val Leu Gly Ala Pro Pro Arg Leu  
                   -10                  -5                  1                  5

Ile Cys Asp Ser Arg Val Leu Glu Arg Tyr Leu Leu Glu Ala Lys Glu  
                   10                  15                  20

Ala Glu Asn Ile Thr Thr Gly Cys Ala Glu His Cys Ser Leu Asn Glu  
                   25                  30                  35

Asn Ile Thr Val Pro Asp Thr Lys Val Asn Phe Tyr Ala Trp Lys Arg  
                   40                  45                  50

Met Glu Val Gly Gln Gln Ala Val Glu Val Trp Gln Gly Leu Ala Leu  
                   55                  60                  65

Leu Ser Glu Ala Val Leu Arg Gly Gln Ala Leu Leu Val Asn Ser Ser  
                   70                  75                  80                  85

Gln Pro Trp Glu Pro Leu Gln Leu His Val Asp Lys Ala Val Ser Gly  
                   90                  95                  100

Leu Arg Ser Leu Thr Thr Leu Leu Arg Ala Leu Gly Ala Gln Lys Glu  
                   105                  110                  115

Ala Ile Ser Pro Pro Asp Ala Ala Ser Ala Ala Pro Leu Arg Thr Ile  
                   120                  125                  130

Thr Ala Asp Thr Phe Arg Lys Leu Phe Arg Val Tyr Ser Asn Phe Leu  
                   135                  140                  145

Arg Gly Lys Leu Lys Leu Tyr Thr Gly Glu Ala Cys Arg Thr Gly Asp  
                   150                  155                  160                  165

Arg

FIG. 2

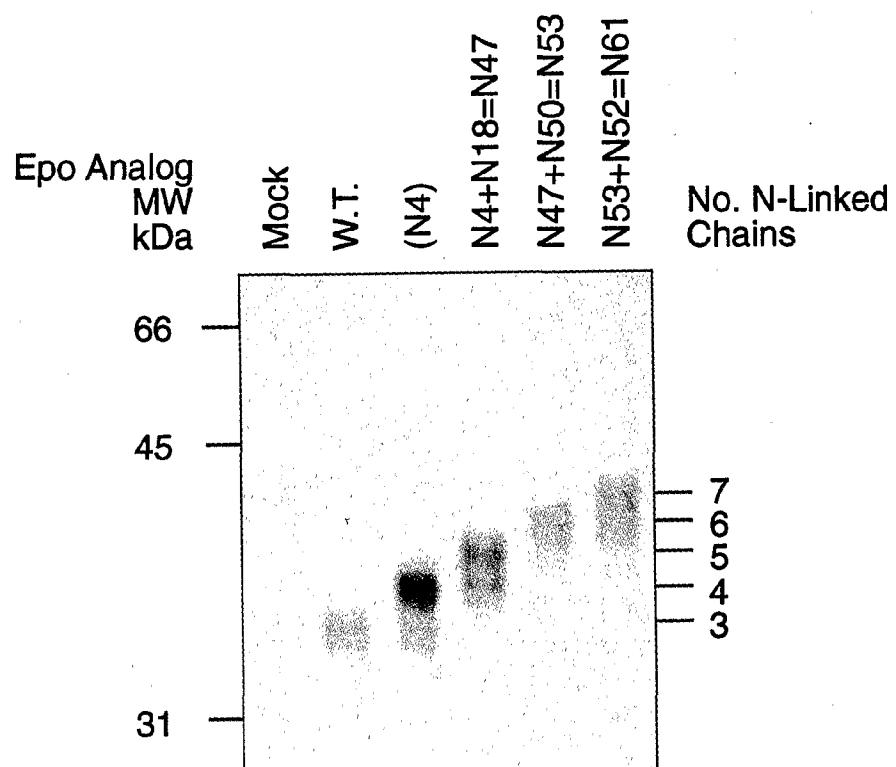


FIG. 3

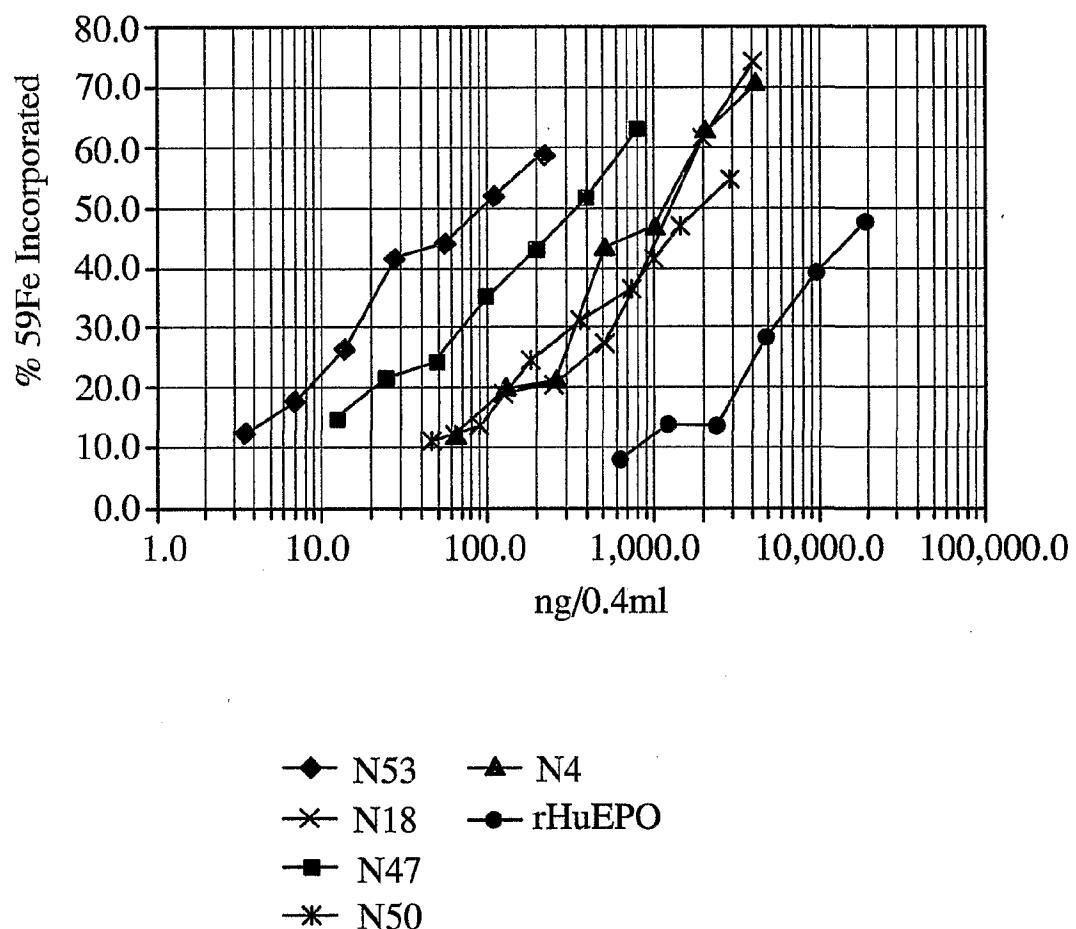


FIG. 4

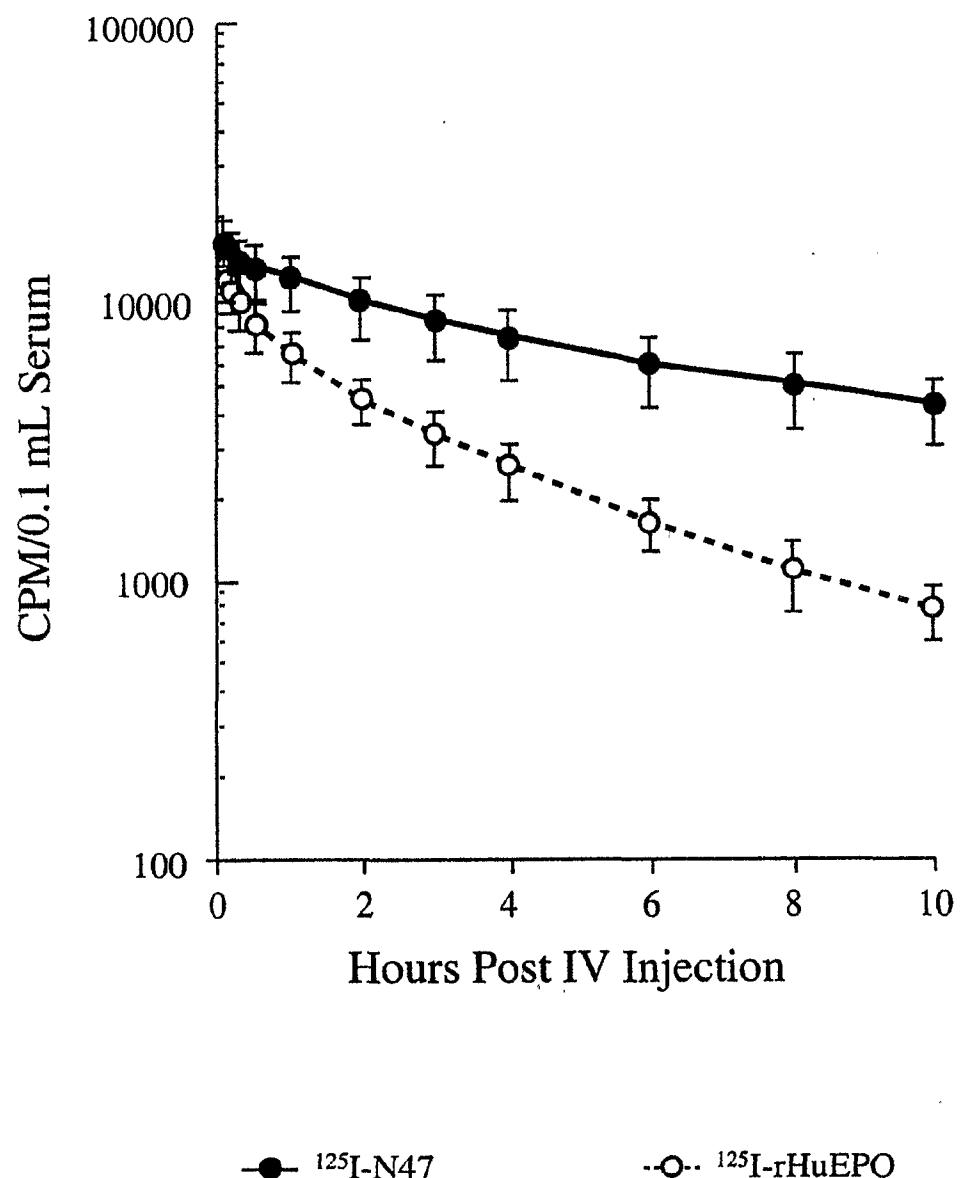


FIG. 5

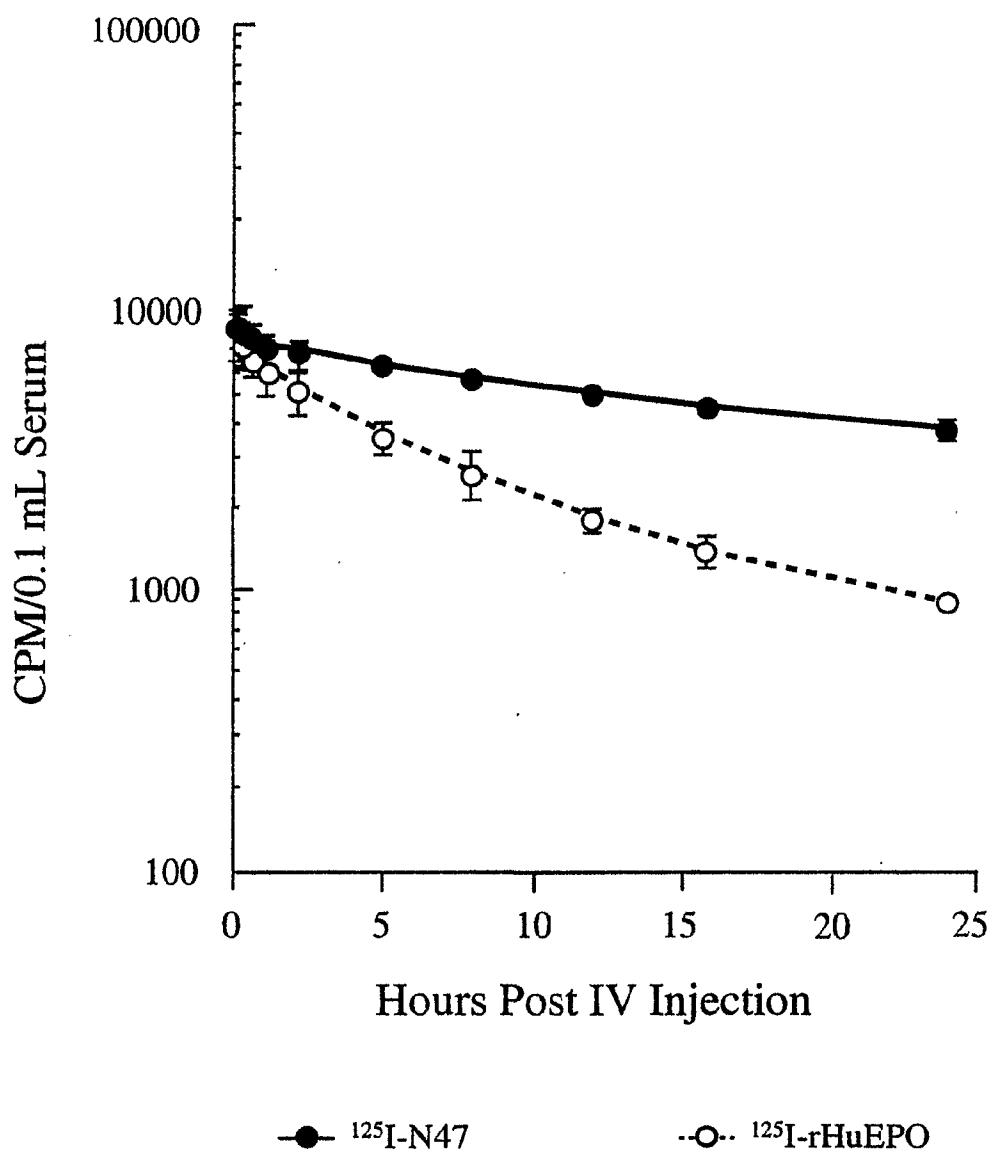


FIG. 6

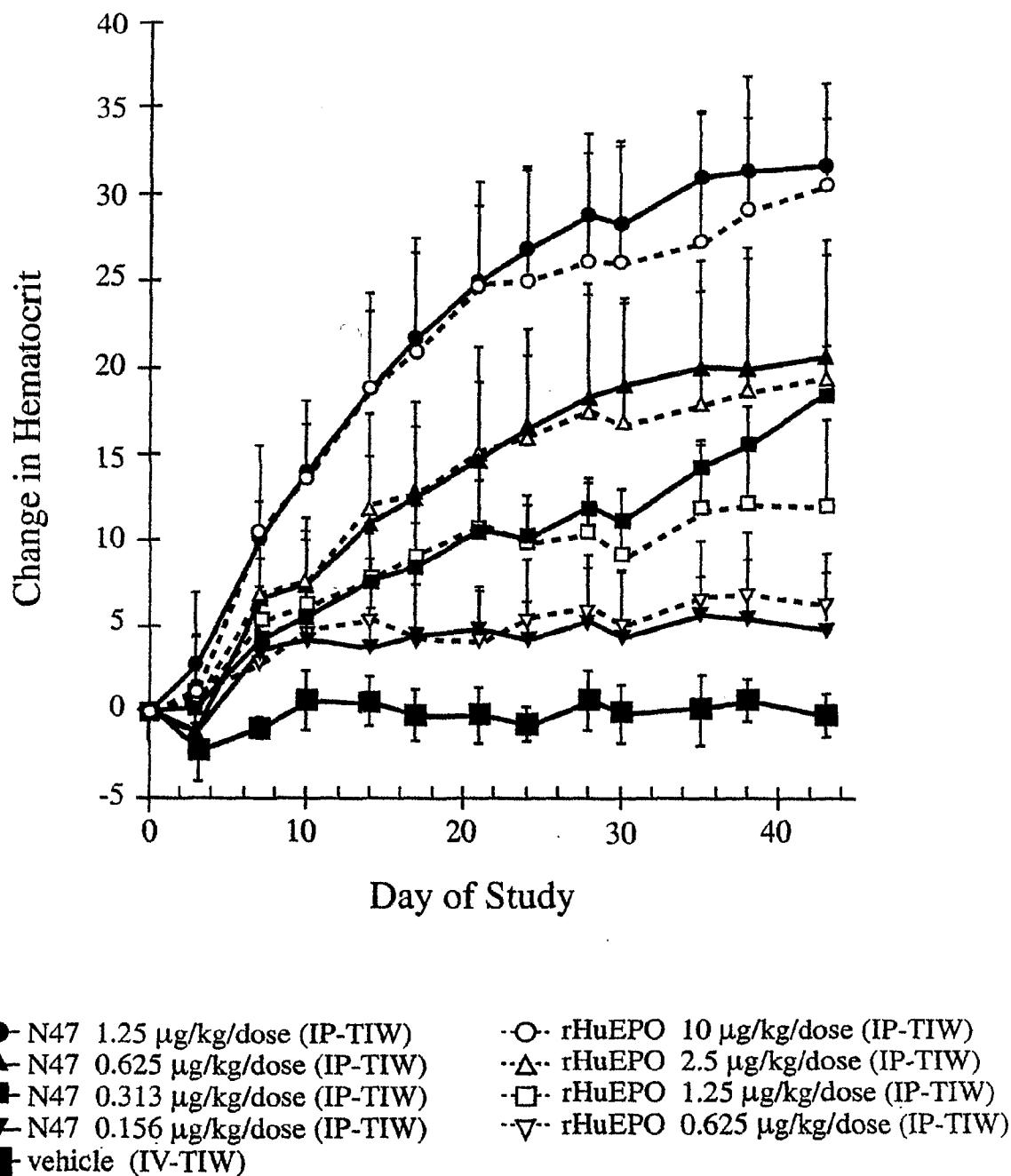


FIG. 7

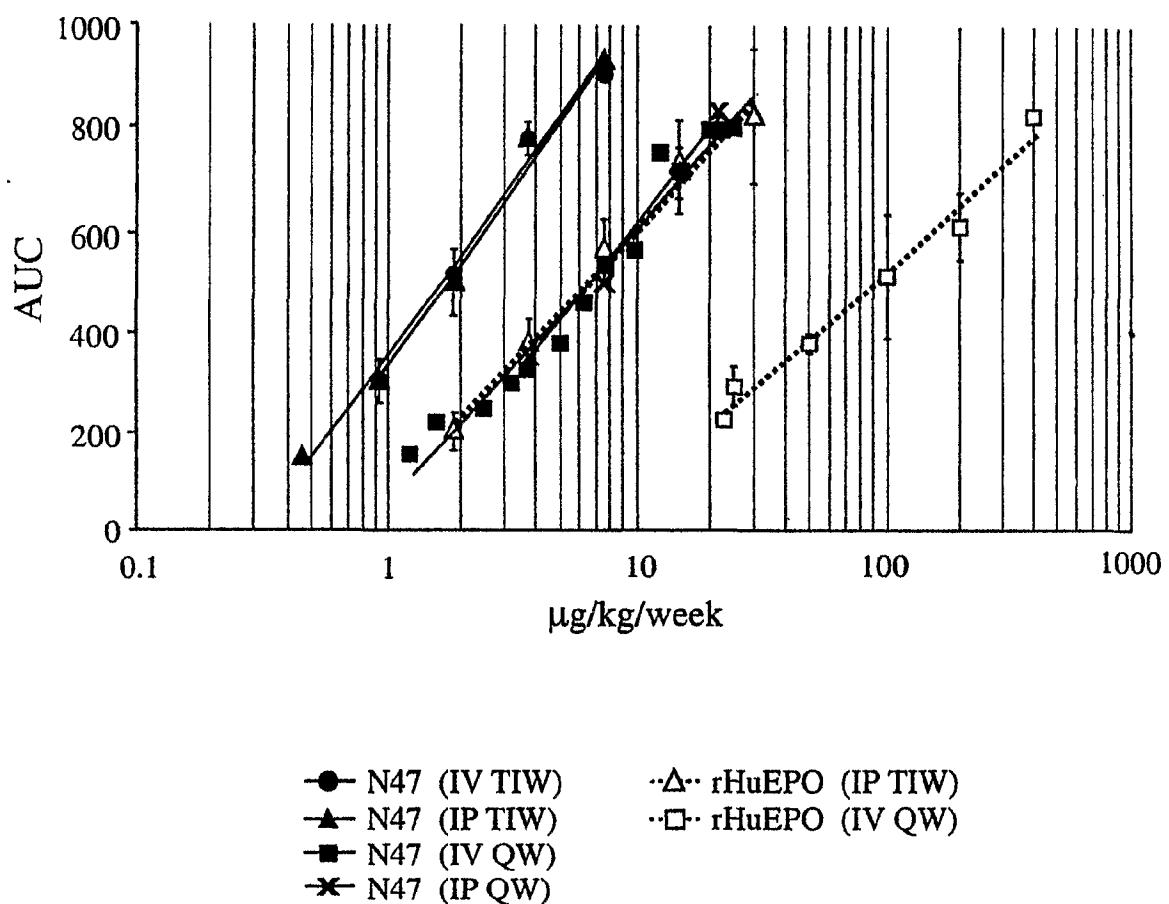


FIG. 8

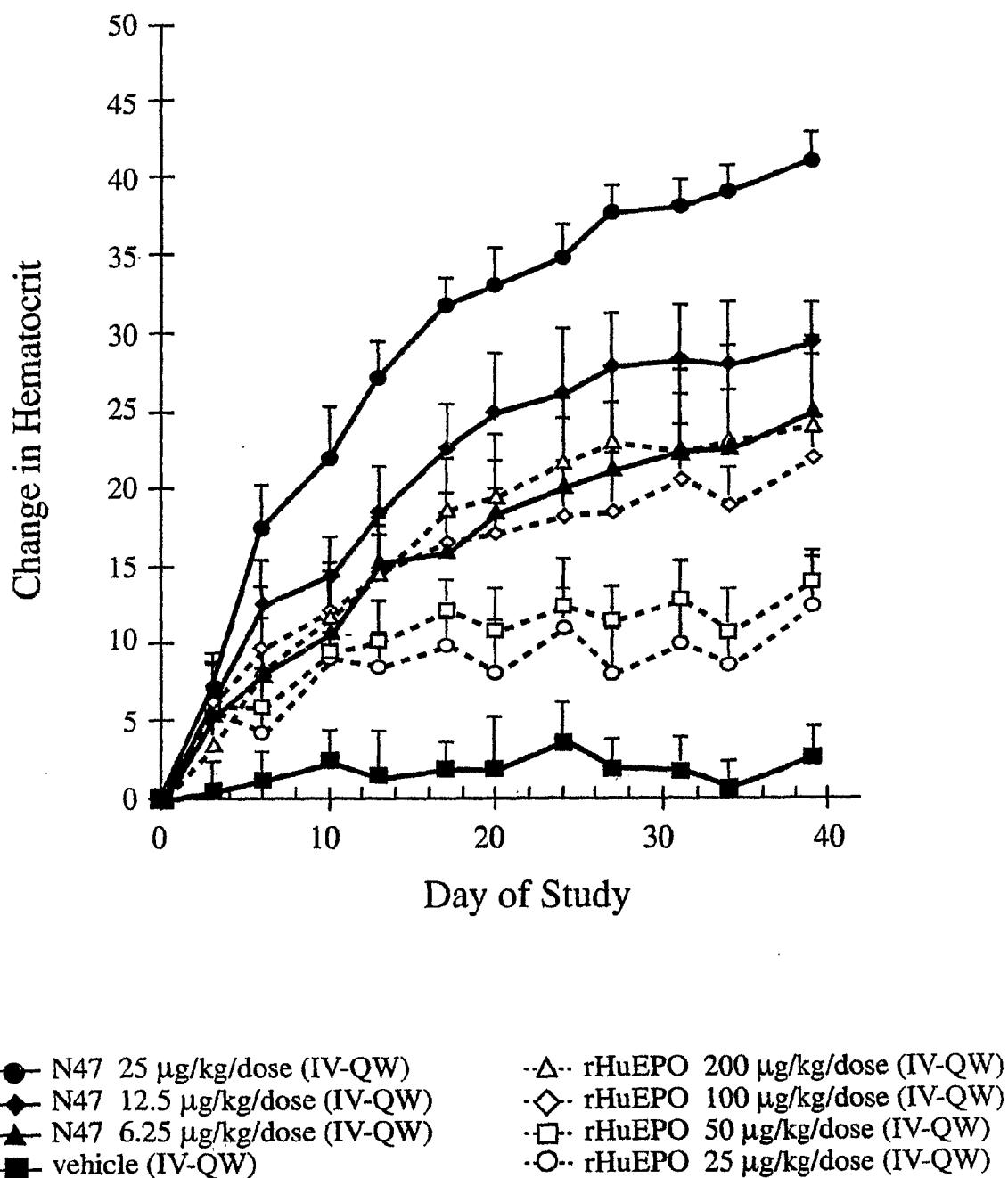
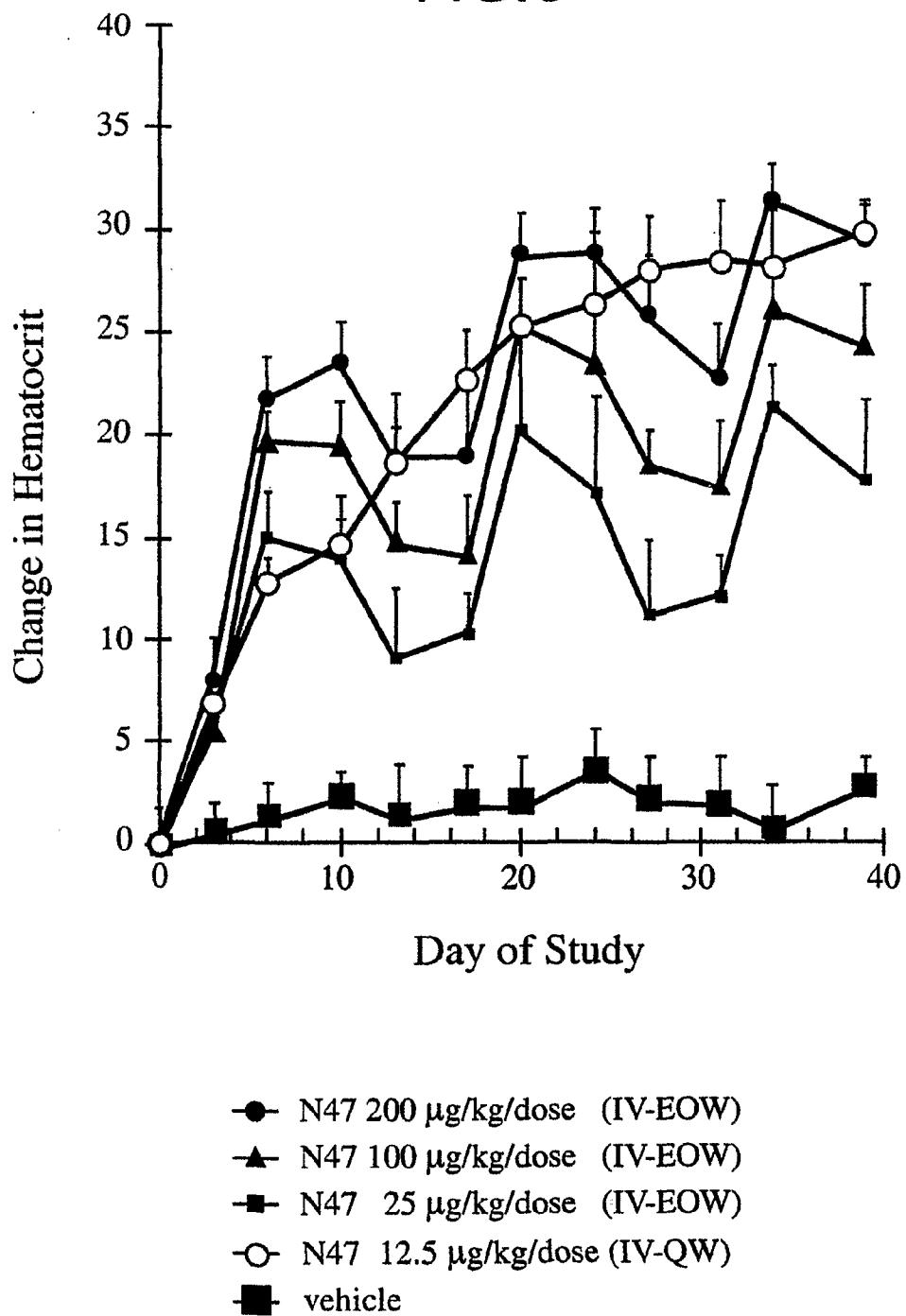


FIG.9



**FIG. 10****Amino acid sequence of hinge, CH2 and CH3 regions human IgG $\gamma$ 1**

Glu	Pro	Lys	Ser	Cys	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro	Cys	Pro	Ala
1				5				10						15	
Pro	Glu	Leu	Leu	Gly	Gly	Pro	Ser	Val	Phe	Leu	Phe	Pro	Pro	Lys	Pro
	20					25						30			
Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr	Cys	Val	Val
	35					40						45			
Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn	Trp	Tyr	Val
	50				55					60					
Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg	Glu	Gln	
	65				70				75			80			
Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val	Leu	His	Gln
	85					90					95				
Asp	Trp	Leu	Asn	Gly	Lys	Glu	Tyr	Lys	Cys	Lys	Val	Ser	Asn	Lys	Ala
	100				105						110				
Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys	Gly	Gln	Pro
	115				120						125				
Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp	Glu	Leu	Thr
	130				135					140					
Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe	Tyr	Pro	Ser
	145				150					155			160		
Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu	Asn	Asn	Tyr
	165				170						175				
Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe	Phe	Leu	Tyr
	180				185						190				
Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly	Asn	Val	Phe
	195				200						205				
Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr	Thr	Gln	Lys
	210				215					220					
Ser	Leu	Ser	Leu	Ser	Pro	Gly	Lys								
	225				230										

## FIG. 11, cont.

661 CTTCCCCCAAAACCAAGGACACCCCTCATGATCTCCGGACCCCTGAGGTACATGCGT  
 661 GAAGGGGGTTTGGTCTGTGGAGTACTAGAGGCCTGGGACTCCAGTGTACGCA 720

b F P P K P K D T L M I S R T P E V T C V -

721 GGTGGTGGACGTGAGCCACGAAGACCCCTGAGGTCAAGTTCAACTGGTACGTGGACGGCGT  
 721 CCACCACCTGCACTCGGTCTCTGGGACTCCAGTTCAAGTTGACCATGCACCTGCGCA 780

b V V D V S H E D P E V K F N W Y V D G V -

781 GGAGGTGCATAATGCCAAGACAAAGCCGCGGGAGGAGCAGTACAACAGCACGTACCGTGT  
 781 CCTCCACGTATTACGGTTCTGTTCGGCCCTCCTCGTCATGTTGTCGTGCATGGCACA 840

b E V H N A K T K P R E E Q Y N S T Y R V -

841 GGTCAAGCGTCCTCACCGTCTGCACCAGGACTGGCTGAATGGCAAGGAGTACAAGTCAA  
 841 CCAGTCGCAGGAGTGGCAGGACGTGGCCTGACCGACTTACCGTCCTCATGTTCACGTT 900

b V S V L T V L H Q D W L N G K E Y K C K -

901 GGTCTCCAACAAAGCCCTCCCAGCCCCATCGAGAAAACCATCTCCAAAGCCAAAGGGCA  
 901 CCAGAGGTTGTTGGGAGGGTAGCTCTTTGGTAGAGGTTCGGTTCCGT 960

b V S N K A L P A P I E K T I S K A K G Q -

961 GCCCCGAGAACACACAGGTGTACACCCCTGCCCATCCGGATGAGCTGACCAAGAACCA  
 961 CGGGGCTTGGTGTCCACATGTGGACGGGGTAGGGCCCTACTCGACTGGTTCTGGT 1020

b P R E P Q V Y T L P P S R D E L T K N Q -

1021 GGTCAAGCCTGACCTGCCTGGTCAAAGGCTTCTATCCCAGCGACATGCCGTGGAGTGGGA  
 1021 CCAGTCGGACTGGACGGACCAGTTCCGAAGATAGGGTCGCTGTAGCGGCACCTCACCT 1080

b V S L T C L V K G F Y P S D I A V E W E -

1081 GAGCAATGGGCAGCCGGAGAACAACTACAAGACCACGCCCTCCGTGCTGGACTCCGACGG  
 1081 CTCGTTACCGTCGGCCTCTGTTGATGTTCTGGTGCAGGGCACGACCTGAGGCTGCC 1140

b S N G Q P E N N Y K T T P P V L D S D G -

1141 CTCCTTCTCCTACAGCAAGCTACCGTGGACAAGAGCAGGTGGCAGCAGGGAACGT  
 1141 GAGGAAGAAGGAGATGTCGTTCGAGTGGCACCTGTTCTCGTCCACCGTCGTCCCCTGCA 1200

S F F L Y S K L T V D K S R W Q Q G N V -

1201 CTTCTCATGCTCCGTGATGCATGAGGCTCTGCACAAACCACTACACGCAGAAGAGCCTCTC  
 1201 GAAGAGTACGAGGCACTACGTACTCCGAGACGTGTTGGTAGTGTGCGTCTCGGAGAG 1260

F S C S V M H E A L H N H Y T Q K S L S -

SalI

1261 CCTGTCTCCGGTAAAtatgtcgac 1286  
 1261 GGACAGAGGCCATTattacagctg

L S P G K \*

## FIG. 11

1 aagcttctagaccaccatgggggtgcacgaatgtcctgcctggctgtggcttcctgtc  
 1 ttcgaagatctggatccccacgtctacaggacggaccacccgaagaggacag 60

M G V H E C P A W L W L L S -

61 cctgctgtcgctccctctggcctccagtcctggcGCCACCACGCCTCATCTGTGA  
 61 ggacacagcgaggagaccggagggtcaggaccgcGGGGTGGTGCAGTAGACACT 120

b L L S L P L G L P V L G A P P R L I C D -  
 121 CAGCCGAGTCCTGGAGAGGTACCTCTGGAGGCCAAGGAGGCCAGAATATCACGACGGG  
 121 GTCGGCTCAGGACCTCTCCATGGAGAACCTCCGGTCCCTCCGGCTTATAGTGCTGCC 180

b S R V L E R Y L L E A K E A E N I T T G -  
 181 CTGtaatGAAacgTGCAGCTTGAATGAGAATATCACTGTCCCAGACACCAAAGTTAATT  
 181 GACAttaCTTtgcACGTCGAACCTACTCTTATAGTGACAGGGTCTGTGGTTCAATTAAA 240

b C N E T C S L N E N I T V P D T K V N F -  
 241 CTATGCCTGGAAGAGGATGGAGGTGGGGCAGCAGGCCGTAGAAGTCTGGCAGGGCCTGGC  
 241 GATACGGACCTCTCCTACCTCCAGCCGTCGTCCGGCATCTTCAGACCGTCCGGACCG 300

b Y A W K R M E V G Q Q A V E V W Q G L A -  
 301 CCTGCTGTGGAAAGCTGTCTGCAGGGGCCAGGCCCTGTTGGTCAACTCTTCCCAGgtgaa  
 301 GGACGACAGCCTTCGACAGGACGCCCGTCCGGACAACCAGTTGAGAAGGGTCactt 360

b L L S E A V L R G Q A L L V N S S Q V N -  
 361 tGAGaCCCTGCAGCTGCATGTGGATAAAGCCGTAGTGGCCTTCGCAGCCTCACCCT  
 361 aCTctGGGACGTCGACGTACACCTATTCGGCAGTCACCGGAAGCGTCGGAGTGGTGA 420

b E T L Q L H V D K A V S G L R S L T T L -  
 421 GCTTCGGGCTCTGGAGGCCAGAAGGAAGCCATCTCCCTCCAGATGCAGGGCCTCAGCTGC  
 421 CGAAGCCCAGACCCCTCGGGTCTCCTCGGTAGAGGGAGGTCTACGCCGGAGTCACG 480

b L R A L G A Q K E A I S P P D A A S A A -  
 481 TCCACTCCGAACAATCACTGCTGACACTTCCGAAACTCTCCGAGTCTACTCCAATT  
 481 AGGTGAGGCTTGTAGTGACGACTGTGAAAGGCCTTGAGAAGGCTCAGATGAGGTAAA 540

b P L R T I T A D T F R K L F R V Y S N F -  
 541 CCTCCGGGAAAGCTGAAGCTGTACACAGGGGAGGCCTGCAGGACAGGGGACAGAGACAA  
 541 GGAGGCCCTTCGACTTCGACATGTGTCCTCGGACGTCCTGTCCCTGTCTCGTT 600

b L R G K L K L Y T G E A C R T G D R D K -  
 601 AACTCACACATGTCCACCTGTCCAGCTCCGAACTCCTGGGGGTCTTCAGTCTCCT  
 601 TTGAGTGTGTACAGGTGGAACAGGTGAGGCCTTGAGGACCCCCCAGGAAGTCAGAAGGA 660

b T H T C P P C P A P E L L G G P S V F L -

## SEQUENCE LISTING

<110> Egrie, Joan  
Elliott, Steven  
Browne, Jeffrey  
Karen, Sitney

<120> Methods and Compositions for the Prevention and Treatment of Anemia

<130> A-460A

<140> 09/559,001  
<141> 2000-04-21

<150> 09/178,292  
<151> 1998-10-23

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<170> PatentIn version 3.0

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20 25 30

Ile Cys Asp Ser Arg Val Leu Glu Arg Tyr Leu Leu Glu Ala Lys Glu  
35 40 45

Ala Glu Asn Ile Thr Thr Gly Cys Ala Glu His Cys Ser Leu Asn Glu  
50 55 60

Asn Ile Thr Val Pro Asp Thr Lys Val Asn Phe Tyr Ala Trp Lys Arg  
65 70 75 80

Met Glu Val Gly Gln Gln Ala Val Glu Val Trp Gln Gly Leu Ala Leu  
85 90 95

Leu Ser Glu Ala Val Leu Arg Gly Gln Ala Leu Leu Val Asn Ser Ser  
100 105 110

Gln Pro Trp Glu Pro Leu Gln Leu His Val Asp Lys Ala Val Ser Gly  
115 120 125

Leu Arg Ser Leu Thr Thr Leu Leu Arg Ala Leu Gly Ala Gln Lys Glu  
130 135 140

Ala Ile Ser Pro Pro Asp Ala Ala Ser Ala Ala Pro Leu Arg Thr Ile  
145 150 155 160

Thr Ala Asp Thr Phe Arg Lys Leu Phe Arg Val Tyr Ser Asn Phe Leu  
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Arg Gly Lys Leu Lys Leu Tyr Thr Gly Glu Ala Cys Arg Thr Gly Asp  
180 185 190

Arg

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35 40 45

Val Asp Val Ser His Glu Asp Pro Glu Val Lys Phe Asn Trp Tyr Val  
50 55 60

Asp Gly Val Glu Val His Asn Ala Lys Thr Lys Pro Arg Glu Glu Gln  
65 70 75 80

Tyr Asn Ser Thr Tyr Arg Val Val Ser Val Leu Thr Val Leu His Gln  
85 90 95

Asp Trp Leu Asn Gly Lys Glu Tyr Lys Cys Lys Val Ser Asn Lys Ala  
100 105 110

Leu Pro Ala Pro Ile Glu Lys Thr Ile Ser Lys Ala Lys Gly Gln Pro  
115 120 125

Arg Glu Pro Gln Val Tyr Thr Leu Pro Pro Ser Arg Asp Glu Leu Thr  
 130 135 140

Lys Asn Gln Val Ser Leu Thr Cys Leu Val Lys Gly Phe Tyr Pro Ser  
 145 150 155 160

Asp Ile Ala Val Glu Trp Glu Ser Asn Gly Gln Pro Glu Asn Asn Tyr  
 165 170 175

Lys Thr Thr Pro Pro Val Leu Asp Ser Asp Gly Ser Phe Phe Leu Tyr  
 180 185 190

Ser Lys Leu Thr Val Asp Lys Ser Arg Trp Gln Gln Gly Asn Val Phe  
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 35 40 45

Gly Cys Thr Thr Cys Thr Cys Cys Thr Gly Thr Cys Thr Thr Cys Gly  
 50 55 60

Ala Ala Gly Ala Thr Cys Thr Gly Gly Thr Gly Gly Thr Ala Cys Cys  
 65 70 75 80

Cys Cys Cys Ala Cys Gly Thr Gly Cys Thr Thr Ala Cys Ala Gly Gly  
 85 90 95

Ala Cys Gly Gly Ala Cys Cys Gly Ala Cys Ala Cys Cys Gly Ala Ala  
 100 105 110

Gly Ala Gly Gly Ala Cys Ala Gly Cys Cys Thr Gly Cys Thr Gly Thr  
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Cys Gly Cys Thr Cys Cys Cys Thr Cys Thr Gly Gly Cys Cys Thr  
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Cys Cys Cys Ala Gly Thr Cys Cys Thr Gly Gly Gly Cys Gly Cys Cys  
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 180 185 190  
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 195 200 205  
 Cys Ala Gly Gly Ala Cys Cys Cys Gly Cys Gly Gly Gly Thr Gly  
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 225 230 235 240  
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 245 250 255  
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 260 265 270  
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 275 280 285  
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 305 310 315 320  
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 340 345 350  
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 355 360 365  
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 420 425 430

Cys Ala Cys Gly Thr Cys Gly Ala Ala Cys Thr Thr Ala Cys Thr Cys  
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Thr Thr Ala Thr Ala Gly Thr Gly Ala Cys Ala Gly Gly Gly Thr Cys  
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Cys Thr Ala Thr Gly Cys Cys Thr Gly Gly Ala Ala Gly Ala Gly Gly  
485 490 495

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515 520 525

Gly Cys Ala Gly Gly Cys Cys Thr Gly Gly Cys Gly Ala Thr Ala  
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Cys Gly Gly Ala Cys Cys Thr Thr Cys Thr Cys Cys Thr Ala Cys Cys  
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Thr Cys Cys Ala Gly Cys Cys Gly Thr Cys Gly Thr Cys Cys Gly  
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Cys Cys Gly Gly Ala Cys Cys Gly Cys Cys Thr Gly Cys Thr Gly Thr  
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Cys Gly Gly Ala Ala Gly Cys Thr Gly Thr Cys Cys Thr Gly Cys Gly  
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Thr Gly Ala Gly Ala Ala Gly Gly Thr Cys Cys Ala Cys Thr Thr  
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Thr Gly Ala Gly Ala Cys Cys Cys Thr Gly Cys Ala Gly Cys Thr Gly

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740	745	750
Thr Cys Ala Gly Thr Gly Gly Cys Cys Thr Thr Cys Gly Cys Ala Gly		
755	760	765
Cys Cys Thr Cys Ala Cys Cys Ala Cys Thr Cys Thr Ala Cys Thr Cys		
770	775	780
Thr Gly Gly Gly Ala Cys Gly Thr Cys Gly Ala Cys Gly Thr Ala Cys		
785	790	795
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805	810	815
Ala Cys Cys Gly Gly Ala Ala Gly Cys Gly Thr Cys Gly Gly Ala Gly		
820	825	830
Thr Gly Gly Thr Gly Ala Gly Ala Gly Cys Thr Thr Cys Gly Gly Gly		
835	840	845
Cys Thr Cys Thr Gly Gly Ala Gly Cys Cys Cys Ala Gly Ala Ala		
850	855	860
Gly Gly Ala Ala Gly Cys Cys Ala Thr Cys Thr Cys Cys Cys Cys Thr		
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Cys Cys Ala Gly Ala Thr Gly Cys Gly Gly Cys Cys Thr Cys Ala Gly		
885	890	895
Cys Thr Gly Cys Cys Gly Ala Ala Gly Cys Cys Cys Gly Ala Gly Ala		
900	905	910
Cys Cys Cys Thr Cys Gly Gly Thr Cys Thr Thr Cys Cys Thr Thr		
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Cys Gly Gly Thr Ala Gly Ala Gly Gly Gly Ala Gly Gly Thr Cys		
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945	950	955
Thr Cys Cys Ala Cys Thr Cys Cys Gly Ala Ala Cys Ala Ala Thr Cys		
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Gly Cys Ala Ala Ala Cys Thr Cys Thr Thr Cys Cys Gly Ala Gly Thr		
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Cys Thr Ala Cys Thr Cys Cys Ala Ala Thr Thr Ala Gly Gly		
1010	1015	1020

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 1490 1495 1500  
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1595	1600	1605
Cys Ala Cys Gly Thr Ala	Cys Cys Gly Thr Gly	Thr Cys Cys Thr
1610	1615	1620
Cys Cys Ala Cys Gly Thr	Ala Thr Thr Ala Cys	Gly Gly Thr Thr
1625	1630	1635
Cys Thr Gly Thr Thr Cys	Gly Gly Cys Gly Cys	Cys Cys Thr
1640	1645	1650
Cys Cys Thr Cys Gly Thr	Cys Ala Thr Gly Thr	Thr Gly Thr Cys
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Gly Thr Gly Cys Ala Thr	Gly Cys Ala Cys Ala	Gly Gly Thr
1670	1675	1680
Cys Ala Gly Cys Gly Thr	Cys Thr Cys Ala Cys	Gly Thr
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Cys Cys Thr Gly Cys Ala	Cys Ala Gly Gly Ala	Cys Thr Gly
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1715	1720	1725
Gly Thr Ala Cys Ala Ala	Gly Thr Gly Cys Ala	Cys Cys Ala
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Gly Thr Cys Gly Cys Ala	Gly Ala Gly Thr Gly	Gly Cys Ala
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Gly Gly Ala Cys Gly Thr	Gly Thr Cys Cys Thr	Gly Ala Cys
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Cys Gly Ala Cys Thr Thr	Ala Cys Gly Thr Thr	Cys Cys Thr
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1790	1795	1800
Cys Thr Cys Cys Ala Ala	Ala Ala Ala Gly Cys	Cys Cys Thr
1805	1810	1815
Cys Cys Cys Ala Gly Cys	Cys Cys Cys Ala Thr	Cys Gly Ala
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Gly Ala Ala Ala Ala Cys	Ala Thr Cys Thr Cys	Cys Ala Ala
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2390	2395	2400												
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2435						2440					2445			
Gly	Ala	Ala	Gly	Ala	Gly	Cys	Cys	Thr	Cys	Thr	Cys	Gly	Ala	Ala
2450						2455					2460			
Gly	Ala	Gly	Thr	Ala	Cys	Gly	Ala	Gly	Gly	Cys	Ala	Cys	Thr	Ala
2465						2470					2475			
Cys	Gly	Thr	Ala	Cys	Thr	Cys	Cys	Gly	Ala	Gly	Ala	Cys	Gly	Thr
2480						2485					2490			
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2495						2500					2505			
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2510						2515					2520			
Gly	Thr	Cys	Thr	Cys	Cys	Gly	Gly	Gly	Thr	Ala	Ala	Ala	Thr	Ala
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2555						2560					2565			
Gly	Cys	Thr	Gly											
				2570										